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<p>(54) Title: NON-STEROIDAL ANTIINFLAMMATORY DRUG FORMULATIONS FOR TOPICAL APPLICATION TO THE SKIN</p> <p>(57) Abstract</p> <p>Topical alcoholic or aqueous alcoholic gels containing ibuprofen or other NSAIDs, such as naproxen, in substantially neutral salt form, have enhanced penetration through skin and may provide rapid pain/inflammation relief by including in the formulation 2-n-nonyl-1,3-dioxolane or other hydrocarbyl derivative of 1,3-dioxolane or 1,3-dioxane or acetal, as skin penetration enhancing compound. The amount of propylene glycol may be varied to adjust the initial flux of the NSAID through the skin, especially for ibuprofen, naproxen, and ketorolac.</p>			

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NON-STEROIDAL ANTIINFLAMMATORY DRUG FORMULATIONS FOR TOPICAL APPLICATION TO THE SKIN

FIELD OF INVENTION

This invention relates to topical compositions for transdermal administration of a non-steroidal antiinflammatory drug (NSAID) through the skin of a patient and to the method for transdermally administering the non-
5 steroidal antiinflammatory drug using the topical composition.

DISCUSSION OF THE PRIOR ART

All drugs must be administered in such a manner that they reach the intended site in the body in an optimal
10 concentration (amount of drug per unit volume of blood) to achieve the desired effect at the proper time, and for an appropriate length of time. Customarily, drugs are taken orally, injected, inhaled, or applied topically. These conventional routes of administration often fail to meet the
15 stated objectives, however. For example, when drugs are absorbed into the blood stream by whatever route, peaks and valleys in the blood concentration of the drug occur and may cause undesirable effects (e.g., peak levels), or loss of activities (e.g., valleys). To meet these problems, a
20 variety of approaches have been investigated. These include, for example, special drug coatings, combining the drug with other materials, suspensions or emulsions, and compressed tablets. Although these formulations attempt to control the release of drugs from their carriers, the

desired effects are often not reproducible, may be subject to patient-to-patient variations, and may not be suitable for prolonged periods of delivery, such as days or even months.

5 The administration of drugs and other biological materials to the bloodstream via a transdermal route of administration has received much attention in recent years. The skin of an average adult covers more than two square meters of surface area and receives about one-third of all
10 blood circulating through the body. It is elastic, rugged, and generally self-generating. The skin consists of three layers: the stratum corneum (S.C.), the epidermis, and the dermis. The stratum corneum represents the rate-limiting step in diffusion of chemical through the skin. The S.C. is
15 composed of dead, keratinized, metabolically inactive cells which are closely packed together, and consists of an amorphous matrix of mainly lipoid and nonfibrous protein within which keratin filaments are distributed. The cells of the S.C. generally contain 20% water, while the cells below, in the stratum germinativum, contain 70% water. The S.C. does not become hydrated readily. Thus, transdermal permeation is primarily controlled by diffusion through the S.C.
20
25

There are several major reasons for the interest in devices for transdermal delivery of drugs:

- elimination of uncertainties of absorption from, and irritation to, the gastrointestinal tract which arise when drugs are administered orally.

- bypassing the portal circulation, thereby eliminating first-pass metabolism in the liver; this is extremely important for drugs with short half-lives, or with potential unwanted actions on the liver.

5 - delivery of medication directly into the systemic circulation at a constant rate (similar to intravenous infusion).

- infrequent dosing (daily, weekly or longer) for certain drugs.

10 - ease of use; foster patient compliance.

However, present transdermal delivery systems have major drawbacks. For example, they are restricted to low-molecular weight drugs and those with structures having the proper lipophilic/hydrophilic balance. High molecular weight drugs or drugs with too high or low hydrophilic balance often cannot be incorporated into current transdermal systems in concentrations high enough to overcome their impermeability through the stratum corneum.

15 Transdermal delivery is generally restricted to those medications requiring delivery rates less than 10 mg/day.

20 In order to obtain higher blood levels, the rate of drug delivery must be increased. There have been many proposals to accomplish the higher rate of drug delivery via the use of absorption promoters and by the development of prodrugs 25 that can be more readily absorbed. Examples of existing absorption enhancers include dimethyl sulfoxide (DMSO), ethylene glycol, hexanol, fatty acid and esters, and

pyrrolidone derivatives, among others. One such enhancer compound which has received much attention is Azone (N-dodecyl azacycloheptan-2-one) developed by Nelson Research Labs., Irvine CA.

5 Certain derivatives of 1,3-dioxanes and 1,3-dioxolanes have previously been used as skin penetration enhancing compounds. These compounds, which have been made commercially available under the trademark SEPA®, are described in detail in U.S. Patent No. 4,861,764. Work with
10 the dioxolane enhancers has been described in several literature and patent publications.

The present inventors have continued to study the effect of the 1,3-dioxane and 1,3-dioxolane derivatives and related acetals as skin penetration enhancer compounds for
15 ibuprofen and other NSAIDs. Surprisingly, it has been found that when propylene glycol is used in the vehicle for the ibuprofen formulations, but not for other NSAIDs, such as diclofenac, ketoprofen, piroxicam, the initial flux rate of ibuprofen decreased as the amount of propylene glycol (PG) increased. Just the opposite effect was observed for the
20 other tested NSAID compounds. In both cases, however, the total payload over a twenty-four hour period is substantially the same. That is, the area under the curve obtained by plotting flux rate over time is the same at 24 hours but the profile of the curves for ibuprofen is dramatically different than for the other tested NSAID compounds.

The present inventors also discovered that at the lower pH's most effective for enhancing the flux of ibuprofen, the 1,3-dioxolane and 1,3-dioxane penetration enhancing compounds become unstable. This problem has now been
5 overcome by incorporating ibuprofen in the form of its substantially neutral (e.g., pH = about 6 to about 8, preferably about 6.5 to 7.5) salt by neutralizing the formulation using an appropriate base, such as sodium hydroxide. This observation of enhancement of the
10 transdermal drug delivery at neutral pH was unexpected since it was originally thought that neutralization of the drug would make it less lipophilic and inhibit its diffusion through the stratum corneum.

It has further been discovered that the flux rates
15 and/or total delivery of NSAIDs, such as, for example, diclofenac, are substantially improved using the 1,3-dioxolane, 1,3-dioxane or corresponding acetal compound skin penetration enhancing compounds.

Another surprising discovery by the present inventors
20 is that for certain of the NSAID active ingredients, such as, naproxen, the permeation through the skin is further enhanced when glycol, e.g., propylene glycol, is omitted from the formulation.

SUMMARY OF INVENTION

25 The present invention has as a principal object to provide stable topical compositions effective for the transdermal application of ibuprofen or other NSAID compounds by the application of the composition to the skin.

The above and other objects of the invention, which will become more apparent from the following more detailed description and preferred embodiments is achieved, according to a first aspect of the invention, by an ibuprofen containing alcoholic or aqueous alcoholic composition which comprises, on a weight basis, of the total composition:

- 5 a therapeutically effective amount of ibuprofen in the form of its pharmacologically acceptable salt;
- 10 a skin penetration enhancing effective amount of a C₇ to C₁₄-hydrocarbyl substituted 1,3-dioxolane, 1,3-dioxane or acetal;
- 15 0 to about 18% of glycol having from 3 to 6 carbon atoms;
- 20 at least 40% of volatile alcohol selected from the group consisting of ethanol, propanol and mixture thereof; 0 to about 40% water; and,
- 25 base to provide a pH in the range of from about 6 to about 8; and optionally, a gelling agent effective to thicken the composition to avoid or minimize run-off when applied to the skin.

In a preferred embodiment of this aspect of the invention the ingredients are included in the formulation in the following ranges:

- 25 from about 2 to 10% ibuprofen;
- from about 2 to 15% of the enhancer wherein the hydrocarbyl group substituent has from about 7 to 10 carbon atoms;

from about 0 to 15% propylene glycol;
from about 55 to 70% ethanol, isopropanol or mixture
thereof;
from about 4 to 35% water; and,
5 base to provide a pH in the range of from about 6.5 to
about 7.5; and
0 to about 2% of cellulosic thickener.

According to another aspect of the invention there are
provided alcoholic or aqueous alcoholic topical compositions
10 effective for the transdermal delivery of non-steroidal
anti-inflammatory drug which comprises, based on the weight
of the total composition,

a therapeutically effective amount of a non-steroidal
antiinflammatory drug selected from the group consisting of
15 heteroaryl acetic acids, arylpropionic acids (other than
ibuprofen), anthranilic acids, enolic acids, alkanones,
sulindac and etodolac;

0.5 to about 25 % of C₇ to C₁₄-hydrocarbyl derivative of
1,3-dioxolane, 1,3-dioxane or acetal as skin penetration
20 enhancer;

0 to about 40% of a glycol having from 3 to 6 carbon
atoms;

at least about 40% of volatile alcohol selected from
the group consisting of ethanol, propanol and mixtures
25 thereof;

0 to about 40% water; and

base to provide a pH of from about 6 to about 8; and
0 to about 5% gelling agent.

According to a preferred embodiment of this second aspect of the invention the composition comprises:

from about 0.1 to 10% diclofenac, ketorolac, naproxen, flurbiprofen, ketoprofen or piroxicam;

5 from about 2 to 15% of the skin penetration enhancer;

0 to about 30% propylene glycol;

from about 35 to 70% ethanol or isopropanol or mixture thereof;

0 to about 20% water;

10 base to provide a pH in the range of from about 6.5 to about 7.5; and

up to about 3% gelling agent.

In still yet another aspect of the invention, the NSAID is naproxen and the glycol component is eliminated from the 15 formulation. According to this third aspect of the invention, there are provided glycol-free topical compositions effective for the transdermal administration of naproxen, which comprise, on a weight basis of the total composition:

20 a pharmaceutically effective amount of pharmacologically acceptable salt of naproxen, from about 2 to 20% of 2-hydrocarbyl group substituted 1,3-dioxolane, 1,3-dioxane, or acetal skin penetration enhancer wherein the hydrocarbyl group has from 7 to 14 25 carbon atoms;

from about 50 to 85% ethanol and/or iso-propanol;
0 to about 40% water; and
base in amount to provide a pH in the range of from
about 6 to about 8; and
5 up to about 5% gelling agent.

BRIEF DESCRIPTION OF THE DRAWINGS

Fig. 1 is a ternary phase diagram showing the miscibility of 2-n-nonyl-1,3-dioxolane skin penetration enhancer at 10 wt.% (■) or 2 wt.% (●) in an ethanol-propylene glycol-water vehicle.
10

Fig. 2 is a ternary phase diagram showing the miscibility of the 1,3-dioxolane skin penetration enhancer at 10 wt.% (●) or 2 wt.% (○) in an isopropanol-propylene glycol-water vehicle.
15

Fig. 3 is a bar graph plotting the flux of ibuprofen Na in an in vitro study as a function of time at 2, 4 or 6 hours post topical application for formulations containing 5% ibuprofen and 0% (bar A), 5% (bar B), 10% (bar C), 15% (bar D) or 20% (bar E) of propylene glycol in an aqueous-alcoholic (ethanol) gel formulation.
20

Fig. 4 is a graph plotting flux of ibuprofen Na versus time in an in vitro study (Example 3) for an aqueous alcoholic gel according to the invention and containing 10 wt.% of 2-n-nonyl-1,3-dioxolane skin penetration enhancer (◊), or a similar gel without skin penetration enhancer (▼), or for four commercial topical ibuprofen preparations: Gelufene® (□), Deep Relief™ (○), Ibutop® (△) and Dolgit® (◆).
25

Fig. 5 is a graph plotting cumulative diffusion of ibuprofen Na versus time for the same samples used in the study of Fig. 4.

Fig. 6 is a graph plotting the number (percentage) of 5 respondents reporting pain relief as a function of time (minutes) in a clinical trial as described in Example 6 using either a formulation according to this invention (O) or a pooled vehicle (□).

Fig. 7 is a graph plotting flux of ibuprofen Na versus 10 time in the *in vitro* study of Example 4 for aqueous alcoholic gels containing 5 wt.% ibuprofen, sodium, 10 wt.% of 2-n-nonyl-1,3-dioxolane skin penetration enhancer and 0% propylene glycol (PG) (▪), 5 wt.% PG (•), 10 wt.% PG (▲), 15 wt.% PG (◊), 20 wt.% PG (◆) or 20 wt.% isopentylidol (IP) 15 (○).

Fig. 8 is a graph plotting flux of ibuprofen Na versus time in the *in vitro* study of Example 5 for aqueous alcoholic gels containing 10 wt.% of penetration enhancer and either 2.5 wt.% ibuprofen, sodium with (▪) or without 20 (◊) propylene glycol (PG); 5 wt.% IB with (•) or without (◆) propylene glycol; and 10 wt.% IB with (▲) or without (○) proplene glycol.

Fig. 9 is a graph plotting flux of ketoprofen versus time using a skin penetration enhancer (SPE) either 25 2-n-nonyl-1,3-dioxolane (SEPA®) at 5% (•) or 10% (▲); diisopropyladipate at 5% (◆) or 10% (□); or benzoate ester at 5% (○) or 10% (△) or without an SPE (▪).

Fig. 10 is a graph plotting cumulative diffusion ($\mu\text{g}/\text{cm}^2$) of ketoprofen versus time for the same samples as used for Fig. 9.

5 Fig. 11 is a graph plotting cumulative diffusion, as percent of dose, versus time for the same samples as used for Fig. 9.

Fig. 12 is a graph plotting flux of ketoprofen versus time using as SPE either 10% 2-n-nonyl-1,3-dioxolane (SEPA®) with a vehicle containing ethylene (E)/propylene glycol (PG)/water (W) at a weight ratio of 70:20:10 (■); or a vehicle containing (E)/glycerol (GL)/(W) (70:20:10) (▲); or a vehicle containing (E)/propylene carbonate (PC)/(W) (70:20:10) (◇); or using 10% decanal as SPE in a (E)/(PG)/(W) (70:20:10) vehicle (●).

15 Fig. 13 is a graph plotting cumulative diffusion ($\mu\text{g}/\text{cm}^2$) versus time for the same samples as used for Fig. 12.

Fig. 14 is a graph plotting cumulative diffusion, as percent of dose, versus time for the same samples as used for Fig. 12.

20 Fig. 15 is a graph plotting flux versus time of ibuprofen from a 5% ibuprofen aqueous gel using as SPE a compound of the invention (●); or cineole (▲) or menthol (■), each in an amount of 10% by weight of the composition.

25 Figs. 16 and 17 are graphs plotting diffusion ($\mu\text{g}/\text{cm}^2$ or percent dose, respectively) versus time for the same samples used for Fig. 14.

DETAILED DESCRIPTION AND PREFERRED EMBODIMENTS

The compositions of the invention are intended for topical, non-invasive, application to the skin, particularly to the region where the non-steroidal anti-inflammatory active ingredient is intended to exert its pharmacological activity, usually to a region of inflammation, injury or pain to the muscles or joints, or other form of cutaneous disorders or disruptions characterized by skin inflammation and/or hyperproliferative activity in the epidermis.

Examples of the non-steroidal antiinflammatory drug (NSAID) which is advantageously administered by the topical formulations of this invention include heteroaryl acetic acids, such as, for example, tolmetin, diclofenac, ketorolac; arylpropionic acids, such as, for example, ibuprofen, naproxen, flurbiprofen, ketoprofen, fenoprofen, oxaprozin; anthranilic acids (fenamates), such as, for example, mefenamic acid, meclofenamic acid, fhenamic acid; enolic acids, such as, for example, oxicams (e.g., piroxicam, tenoxicam), pyrazolidinediones (e.g., phenylbutazone, oxyphenthatrazone); alkanoes, such as, for example, nabumetone. Among these, especially preferred, based on the current level of knowledge in the pharmacological arts, are ibuprofen, diclofenac, ketorolac, naproxen, flurbiprofen, ketoprofen and piroxicam. More generally, however, any of the government approved NSAIDs, such as listed in, for example, the most current edition of The Merck Index, may be advantageously used.

According to the present invention the NSAID is administered in the form of its pharmacologically acceptable substantially neutral salt. The formulations are made substantially neutral by addition of a pH modifying agent 5 (base) in an amount to provide a pH in the range of from 6.0 to 8.0, preferably from 6.5 to 7.5, especially preferably from 6.8 to 7.4, such as 7.0. Any of the well known and pharmacologically safe inorganic or organic basic compounds can be used for this purpose and examples include 10 inorganic salt, such as the sodium or other alkali or alkaline earth metal salts such as hydroxides, e.g., sodium hydroxide or potassium hydroxide; ammonium salt; or organic salt, especially amine salt, such as, for example, diethylamine; diethanolamine, triethanolamine, 15 diisopropanolamine, N-methylglucamine, ethanolamine, isopropylamine, tetrahydroxypropyl ethylene diamine methylamine, ethylamine, propylamine, and the like.

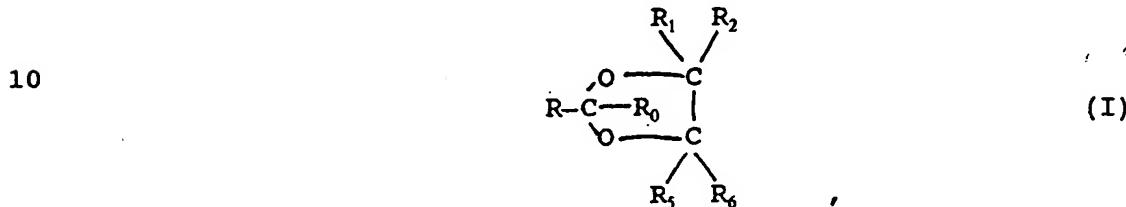
For any particular formulation the NSAID and other ingredients may be selected to achieve the desired drug 20 delivery profile and the amount of penetration desired. The optimum pH may then be determined and will depend on, for example, the nature of the NSAID, the base, and degree of flux required.

The penetration of the active ingredient through the 25 skin is enhanced to an acceptable level by including in the composition a skin penetration enhancing effective amount of

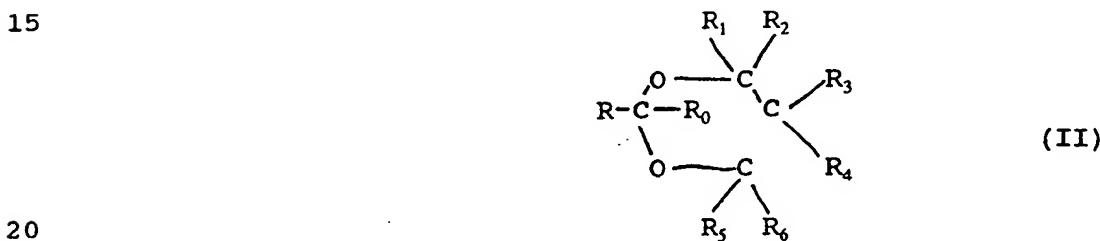
an enhancer compound of the substituted 1,3-dioxacyclopentane and substituted 1,3-dioxacyclohexane types disclosed in U.S. 4,861,764, the disclosure of which is incorporated herein in its entirety by reference thereto, or 5 the corresponding substituted acetal compound.

Representative examples of the skin penetration enhancing compounds include:

2-substituted 1,3-dioxolanes of the formula (I):



2-substituted 1,3-dioxanes of the formula (II):



substituted-acetals of the formula (III):



In the above formulas (I), (II) and (III) R preferably represents a C₁ to C₁₄ hydrocarbyl group,

30 R₀, R₁, R₂, R₃, R₄, R₅, and R₆, each, independently, represent hydrogen or a C₁ to C₄ alkyl group.

R₁' and R₂', each, independently, represent C₁ to C₄ alkyl group.

The hydrocarbyl group for R may be a straight or branched chain alkyl, alkenyl or alkynyl group, especially alkyl or alkenyl. Preferably, R represents a C₁ to C₁₂ aliphatic group; especially C₁ to C₁₀ aliphatic group.
5 Examples of suitable alkyl groups include, for example, n-hexyl, n-heptyl, n-octyl, n-nonyl, n-decyl, n-undecyl, n-dodecyl, 2-methyl-octyl, 4-ethyl-decyl,
10 8-methyl-decyl, and the like. The straight chain alkyl groups, such as n-heptyl, n-octyl, n-nonyl and n-decyl, are especially preferred. Examples of alkenyl groups include, for example, 2-hexenyl, 2-heptenyl, 2-octenyl, 2-nonenyl, 2',6'-dimethyl-2',6'-heptadienyl, 2'6'-dimethyl-2'heptaenyl,
15 and the like. The R group may also be substituted by, for example, halo, hydroxy, carboxy, carboxamide and carboalkoxy.

The C₁ to C₄ alkyl group may be, for example, methyl, ethyl, n-propyl, isopropyl, n-butyl, tert-butyl, and the like. The preferred alkyl groups for R₀, and for R₁ to R₆ and for R₁' and R₂', are alkyl having 1 or 2 carbon atoms, most especially ethyl. R₀, and R₁ to R₆ may also, preferably, all be hydrogen.
20

Specific enhancer compounds include, for example, 2-n-pentyl-1,3-dioxolane, 2-n-heptyl-1,3-dioxolane, 2-n-nonyl-1,3-dioxolane, 2-n-undecyl-1,3-dioxolane, 2-n-nonyl-

1,3-dioxane, 2-n-undecyl-1,3-dioxane, 2-n-heptylaldehyde-acetal, 2-n-octyl-aldehyde-acetal, 2-n-nonylaldehyde-acetal, 2-n-decylaldehyde-acetal, 3,7-dimethyl-2,6-octadienal (citral), citronal and the like. 2-n-nonyl-1,3-dioxolane is especially preferred and is commercially available from MacroChem Corporation of Lexington, Massachusetts, under the trademark SEPA®.

The amount of the enhancer compound is selected to provide the desired delivery rate for the active compound but, taking into consideration such additional factors as, product stability, side effects, carrier system and the like. Generally, depending on the particular NSAID and other vehicles, amounts in the range of from about 0.5 to 25%, preferably from about 2 or 3 to 12 or 15 percent, especially from about 5 to 10 percent, of the composition, will provide optimal flux rate and 24 hour payload of the active ingredient. Usually, for cream formulations the amount of enhancer compound may be lower than for gel formulations, such as from about 2 to 10 percent of the formulation.

The compositions are generally formulated as gels, especially aqueous-alcoholic gels. However, other forms, such as, for example, lotions, creams, mousses, aerosols, ointments, lubricants, etc., may be used so long as when applied to the affected area of the skin the formulation will stay in place, i.e., without run-off, for sufficient time, to permit an individual to spread and retain the composition over and on the affected area.

The vehicle for any of the forms of the compositions of the invention will include glycol, e.g., propylene glycol, butylene glycol, hexylene glycol, etc. (except in the case of the third embodiment described above), lower 5 alcohol, e.g., ethanol, isopropanol, and, usually, water. A thickening or gelling agent is also usually and preferably included to facilitate application of the formulation to the skin. In addition, of course, the skin penetration enhancing dioxolane, dioxane or acetal is included in the 10 formulations in an amount effective to enhance the penetration of the active NSAID ingredient through the skin, including the stratum corneum.

Accordingly, the vehicle or carrier system for the NSAID and enhancer components is preferably an aqueous or 15 non-aqueous alcoholic carrier containing sufficient alcohol, especially ethanol and/or isopropanol and, often, glycol, e.g., propylene glycol, to solubilize the NSAID and be miscible with the enhancer. Generally, however, depending on the amounts of enhancer and NSAID in the formulations 20 the aqueous alcoholic carrier can contain from about 35% to about 70% of ethyl alcohol and/or isopropyl alcohol, preferably, from about 50 to about 70 percent of ethanol or from about 45 to 55 percent of isopropanol. Mixtures of ethanol and isopropanol in proportions providing the desired 25 solubility of NSAID and compatibility with the enhancer can also be used. More generally, however, the present

inventors have developed miscibility data for combinations of alcohol (ethanol or isopropanol), glycol (propylene glycol) and water for the enhancer (2-n-nonyl 1,3-dioxolane). This data is graphically represented by the 5 ternary phase diagrams provided as Figure 1 (for ethanol) at 2 wt.% (•) and 10 wt.% (■) of the enhancer compound and Figure 2 (for isopropanol) at 2 wt.% (○) and 10 wt.% (●) of the enhancer compound. In each of these phase diagrams, the upper portions (above the lines connecting the data points) represent the proportions at which the vehicle components 10 are miscible with each other and with the enhancer; conversely, the region below the lines connecting the data points represent the proportions where the vehicle components are immiscible.

15 Again, the total amount of the aqueous or non-aqueous, alcoholic carrier will depend on the amount of NSAID, amount and type of enhancer, and the form of the composition, e.g., gel, cream, ointment, etc. Usually amounts of the aqueous or non-aqueous alcoholic carrier within the range of from 20 about 70% to about 95% may be used.

In the preferred compositions which are in the form of a gel, a thickening agent, such as hydroxypropyl cellulose, will be included as a gelling agent. However, any other pharmaceutically acceptable thickening/gelling 25 agent may be used. For example, mention may be made of other cellulosic ethers, polymeric thickening agents, e.g.,

acrylic acid polymers, Carbopol® thickeners, etc., xanthan gum, guar gum, and the like, as well as inorganic thickeners/gelling agents. The amount of the thickening agent is not particularly critical and can be selected to provide the desired product consistency or viscosity to allow for easy application to the skin but which will not be too watery or loose so that it will stay where applied. Generally, depending on its molecular weight, amounts of thickening agent up to about 5%, such as, for example, from 0.1 to about 2%, of the composition will provide the desired effect.

As also well known in this art, it is possible to include other ingredients in the formulations for particular aesthetic and/or functional effects. For example, the formulations may, optionally, include one or more moisturizers for hydrating the skin and emollients for softening and smoothing the skin. Glycerin is an example of such a suitable moisturizing additive. When present the additive will usually be incorporated in an amount of up to about 5 percent by weight of the composition, for example, from about 0.1 to 5%.

The effects of the topical compositions according to the invention are further illustrated by way of the following representative examples which in no way are intended to limit the scope of the invention.

Example 1

This example compares the percutaneous absorption through porcine skin, of ibuprofen from aqueous alcoholic gels containing 5 wt.% ibuprofen and either 5%, 10% or 15% of 2-n-nonyl-1,3-dioxolane, using an ethanol/water carrier at a 70:30 mixing ratio. The formulations include NaOH to adjust the pH to 7.4, but do not include a glycol.

Hydroxypropyl cellulose (2 wt.%) is used as the gelling agent. The test compositions are applied to provide about 10 30 milligrams (mg) of the composition per square centimeter (cm^2) of porcine skin.

The tests are run in standard static cells with phosphate buffered saline (PBS) as the receptor fluid (surface area 0.635 cm^2 , temperature 32°C). The following 15 Table 1 shows the total amount of ibuprofen applied to the skin for each formulation. The differences result from the slightly different thicknesses at which the test formulations are applied. Each test was run for 24 hours under non-occluded conditions with the finite dose of the 20 test formulation.

Table 1

Enhancer	Amount of Enhancer (wt.%)	Total Amount of Ibuprofen applied to skin sample (μ g) per 0.635 cm^2 cell
2-n-nonyl- 1,3-dioxolane	5	988
2-n-nonyl- 1,3-dioxolane	10	1051
2-n-nonyl- 1,3-dioxolane	15	1038

The results are obtained and reported in Table 2 as the average values for six (6) cells (samples). The initial flux over the first two hours was significantly higher for each formulation containing the 1,3-dioxolane, but especially at the higher level (15%, maximum flux about 8.5 $\mu\text{g}/\text{cm}^2/\text{hr}$; 5% and 10%, maximum flux about 4 $\mu\text{g}/\text{cm}^2/\text{hr}$) of the enhancer. The flux tended to even out after 4 to 6 hours and continued at about the same level for at least about 24 hours. The results for total amount of ibuprofen vs. time (flux); and the payout of ibuprofen at 24 hours, total and percent of dose, are shown in the following Table 2:

Table 2

Amount Enhancer (%)	2h	4h	6h	24h	Delivery at 24h	
					Total (μg)	% of Dose
15	5	4	2	1.5	1	20.2 \pm 6.9 2.1 \pm .7
	10	4	2	1.5	1.5	26.6 \pm 8.1 2.4 \pm .6
	15	8	4	3.5	3.5	57.7 \pm 33.4 5.4 \pm 2.5

When the procedure of Example 1 was repeated but using 5%, 10% or 15% of Azone™, the initial flux at 2 hours and 4 hours was only between about 1 to 1.5 µg/cm²/hr.

Example 2

5 This example shows the effect of incorporating propylene glycol in the aqueous alcoholic gel formulation containing 5% ibuprofen and 10% 2-n-nonyl-1,3-dioxolane using an ethanol:water vehicle at a 70:30 weight mixing ratio. The compositions used in these tests are shown in
10 Table 3 (NaOH is added to adjust the pH to 7.4):

Table 3

	ibuprofen (%)	enhancer (%)	propylene glycol (%)	Ethanol (%)	Water (%)	Total (%)
A	5	10	0	59.5	25.5	100
B	5	10	5	56	24	100
C	5	10	15	52.5	22.5	100
D	5	10	15	49	21	100
E	5	10	20	45.5	19.5	100

The test was run using the same conditions as described in Example 1. The flux was measured at 2, 4 and 6 hours. The results are shown graphically in Fig. 3. From this figure it is seen that the flux at 2 hours decreases nearly linearly as the propylene glycol (PG) content increases from 0% to 5% to 10% to 15% to 20%. At four hours after the composition is applied to the test skin sample the fluxes for each concentration of PG has increased but more so for the compositions containing the higher amounts of PG.
15 Finally, at 6 hours the fluxes begin to even out.
20

This example, therefore, shows that only low or no propylene glycol should be included in the ibuprofen topical composition using the 2-substituted-1,3-dioxolane, 2-substituted-1,3-dioxane or substituted acetal as the penetration enhancer where the goal is to administer large quantity of active ingredient as quickly as possible such that relief from pain or inflammation can begin rapidly, for example in the treatment of sunburn or other burn injury or for relief of muscular pain caused by inflammation. However, over the longer period of time essentially the same amount of ibuprofen is percutaneously delivered at each amount of propylene glycol.

Example 3

This example is similar to Example 1 but compares a topical aqueous alcoholic gel formulation with ibuprofen according to the present invention with a similar gel but without the enhancer and with four other commercially available topical ibuprofen formulations. Also, human skin was used rather than porcine skin. The composition according to the present invention and the comparison were as follows:

<u>Ingredient</u>	<u>Invention Amount (wt. %)</u>	<u>Comparison Amount (wt. %)</u>
Ibuprofen	5	5
2-n-nonyl-1,3-dioxolane	10	0
Ethanol	59	65
Propylene glycol	17	19
Water	7	9

Hydroxypropyl

cellulose

2

2

Sodium Hydroxide q.s. to pH 7 q.s. to pH 7

The commercially available products were: Gelufene® (ibuprofen 5%, isopropyl alcohol, hydroxyethylcellulose, sodium hydroxide, benzyl alcohol and purified water), Dolgit® cream (ibuprofen 5%, medium chain triglycerides, mixture of glycerol monostearate and polyoxyethylene stearates, polyoxyethylene fatty acid esters, xanthan gum, lavender oil, neroli oil, water, propylene glycol, parahydroxybenzoate of methyl soda), Ibutop® (ibuprofen 5%) (Laboratoire Chefaro-Ardeval, Saint-Denis Cedex, France) and Deep Relief™ gel (ibuprofen 5%, menthol, Carbomer, propylene glycol, di-isopropanolamine, ethanol, purified water). The results are shown in Figure 4 for flux versus time and in Figure 5 for cumulative diffusion of ibuprofen through the skin sample. The initial flux and cumulative amount of the ibuprofen are both significantly higher for the invention formulation than for the control or commercial products.

Example 4

This Example is also similar to Example 1 but is designed to compare the percutaneous absorption through porcine skin of ibuprofen (as the sodium salt) from 5% ibuprofen formulations containing 10% 2-n-nonyl-1,3-dioxolane with 5%, 10%, 15% or 20% propylene glycol (PG) or 20% isopentyldiol (IP). The test conditions were otherwise the same as used in Example 1. The formulations tested are shown in Table 4:

Table 4

Run No.	Ibuprofen (%)	Enhancer (%)	glycol PG or IP (%)	EtOH (%)	Water (%)	Total (%)
1	5	10	0 PG	59.5	25.5	100
2	5	10	5 PG	58.9	21.1	100
3	5	10	10 PG	58.3	16.7	100
4	5	10	15 PG	57.6	12.4	100
5	5	10	20 PG	56.9	8.1	100
6	5	10	20 IP	56.9	8.1	100

The flux of ibuprofen in the receptor cell was measured and the results are shown in Fig. 7. In this case, the peak flux was reached for the formulations of Run Nos. 1-4 within four hours after application and reached values in the range of from about 60 to 72 $\mu\text{g}/\text{cm}^2/\text{hr}$. For the 20% PG formulation (run #5) the peak flux was reached at 6 hours and was about 50 $\mu\text{g}/\text{cm}^2/\text{hr}$. For the 20% IP formulation (run #6) the peak flux was only about 16 $\mu\text{g}/\text{cm}^2/\text{hr}$ and was not reached until 8 hours after application.

10 The following Table 5 shows the results reported for the cumulative amount of ibuprofen reaching the receptor cell after 24 hours, and the amount reaching the receptor cell as a percentage (%) of the dose applied after 24 hours.

Table 5

15

Delivery at 24h

Run	<u>PG or IP (wt.%)</u>	<u>Total (μg)</u>	<u>% Dose</u>
1	0	541.7 ± 4.8	60.1 ± 8.6
2	5 PG	519.8 ± 53.7	57.9 ± 5.7
3	10 PG	498.6 ± 58.9	56.5 ± 6.6
20	4 PG	443.5 ± 105.1	50.1 ± 11.4

5	20 PG	358.5 ± 73.5	40.7 ± 8.4
6	20 IP	144.5 ± 31.1	16.6 ± 3.6

Example 5

This example is designed to demonstrate the effect of concentration of ibuprofen (IB) on flux and total delivery (24h) for formulations with and without propylene glycol. The test were run under the same conditions as in Example 1 except that human skin was used, an 80/20 mixture of PBS and ethanol was used as the receptor fluid, and the pH was adjusted to 7.7 with sodium hydroxide; the test compositions which were prepared and tested (the enhancer was 2-n-nonyl-1,3-dioxolane) are shown in the following Table 6:

Table 6

Run No.	Ibuprofen (%)	Enhancer (%)	PG (%)	EtOH (%)	Water (%)	Total (%)
1	2.5	10	17.5	61.25	8.75	100
2	5	10	17	59.5	8.5	100
3	10	10	16	56	8	100
4	2.5	10	-	61.25	26.25	100
5	5	10	-	59.5	25.5	100
6	10	10	-	56	24	100

The flux as a function of time for each of the test formulations is shown in Fig. 8. Comparing the results for Run Nos. 1 and 4, Run Nos. 2 and 5 and Run Nos. 3 and 6, it is seen that in each case the maximum (peak) flux and time to reach peak flux were higher and quicker for the formulations without propylene glycol.

The results for the cumulative dose of ibuprofen at 24 hours and the percentage of the original dose passing through the skin at 24 hours are shown in the following

Table 7:

5

Table 7

Delivery at 24h

Run	<u>PG(wt%)/IB(wt%)</u>	<u>Total (μg)</u>	<u>% Dose</u>
1	17.5/2.5	193.5 ± 37.3	37.6 ± 6.5
2	17/5	361.9 ± 116.3	38.2 ± 12.7
10	16/10	615.2 ± 152.6	29.8 ± 7.5
	4/2.5	392.2 ± 189.1	80.4 ± 30.2
	0/5	535.7 ± 189.1	53.6 ± 19.9
	0/10	805.9 ± 214.4	41.2 ± 8.8

These results show, for example, that the formulation with 10% ibuprofen but without propylene glycol (Run No. 6) gives the highest flux. Furthermore, the formulation of Run No. 4 (2.5g IB per 100 g formula + 0 PG) will deliver 80 mg of ibuprofen over a 100 cm² area.

Example 6

This example shows the results of a clinical study on human patients experiencing severe pain. The same formulation as shown in Example 3 (10% 2-nonyl-1,3-dioxolane and 5% ibuprofen and 17% propylene glycol) is used in these studies. As control, a pooled vehicle (mixture of first formulation with no drug and no enhancer and second formulation with no drug but with enhancer) was similarly tested. The results are shown in Fig. 6.

Example 7

This Example compares percutaneous absorption through human skin of Naproxen Na from various formulations containing 5% (w/w) Naproxen gels with or without propylene glycol and with or without skin penetration enhancer compound (2-n-nonyl-1,3-dioxolane). The tests were run under the same conditions described in Example 5 (pH = 7.7) using the gel formulations as shown in Table 8:

Table 8

Run No.	Naproxen (%)	Enhancer (%)	PG (%)	EtOH (%)	Water (%)	Total (%)
1	5	0	-	66.5	28.5	100
2	5	5	-	63	27	100
3	5	10	-	59.5	25.5	100
4	5	0	19	66.5	9.5	100
5	5	5	18	63	9	100
6	5	10	17	59.5	8.5	100

With these Naproxen gel formulations the highest flux and highest total delivery was achieved with the formulation of Run No. 2 (0% PG + 5% enhancer). The peak flux was observed 4 hours after application of the gel on the skin.

The results for the cumulative dose of naproxen at 24 hours and the percentage of the original dose passing through the skin at 24 hours are shown in the following

15 Table 9:

Table 9

Delivery of Naproxen Na at 24h

Run	<u>PG(wt%) / enhancer(wt%)</u>	<u>Total (μg)</u>	<u>% Dose</u>
1	0/0	51.5 ± 16.6	5.2 ± 1.4
5	2	499.0 ± 96.9	52.3 ± 10.9
	3	369.1 ± 74.8	35.9 ± 7.5
	4	29.4 ± 12.4	2.9 ± 1.1
	5	149.5 ± 40.3	15.0 ± 4.0
	6	409.6 ± 113.8	39.7 ± 12.3

10 These results appear to show that propylene glycol is not functioning as an enhancer in the subject formulations (compare, e.g., Run No. 1 with Run No. 4) and further, propylene glycol has an adverse impact on the delivery of the NSAID, Naproxen.

15 Example 8

This example illustrates the effect of propylene glycol (PG) on delivery of various NSAIDs from aqueous formulations containing 10 wt.% of skin penetration enhancer, 2-n-nonyl-1,3-dioxolane. All the tested formulations included ethanol and water at a 70:30 weight ratio and were neutralized with base to a pH of about 7. The tests were run in standard static cells under substantially the same conditions as described in Example 1 but using human skin rather than procine skin. The tested compositions and results are shown 20 in the following Table 10.

25

Table 10

NSAID Drug	Drug conc.	% enhancer	% PG	Avg 24 Hour % Dose
Ketoprofen	2.5%	10	0	9.1
		10	5	10.3
		10	10	12.0
		10	20	27.0
Piroxicam	0.5%	10	0	66.3
		10	10	55.1
		10	20	63.5
Ibuprofen	5%	10	0	61.7
		10	5	57.9
		10	10	56.5
		10	15	50.1
		10	20	28.3
Diclofenac	1%	10	0	18.7
		10	5	29.7
		10	10	35.3
		10	20	45.2
Ketorolac	1%	10	0	10.8
		10	5	27.1
		10	10	25.0
		10	20	19.3
Naproxen	5%	10	0	35.9
		10	20	39.7

As may be readily ascertained from these results the effects of propylene glycol differs substantially from one NSAID to another. The effect, in terms of total delivery (reported as percent of original dose) at twenty four hours) of NSAID is positive for ketoprofen and diclofenac; substantially neutral for piroxican and naproxen; negative for ibuprofen (particularly at the high levels of propylene glycol); and intermediate for ketorolac.

5

Example 9

This example further illustrates the effects of PG on drug delivery (0.5% piroxicam) at two different levels of the enhancer, 2-n-nonyl-1,3-dioxolane (5% or 10%) versus a control (0% enhancer, 0% PG) and a commercial product, Geldene® (0.5% piroxicam in the form of its diisopropanolamine (DIPA) salt; approximately 24% ethanol; >0 PG). In the compositions according to the invention and the control triethanolamine (TEA) was used as the base to neutralize the piroxicam and the vehicle was ethanol:water (70:30). The formulations and test procedures were, otherwise, as described in Example 8. The results are shown below in Table 11.

Table 11

15	Run No.	Propylene Glycol (wt%)	Enhancer (wt.%)	Peak Flux $\mu\text{g}/\text{cm}^2/\text{h}$	% of Dose 24h
20	9-1	0	10	8.7	66
	9-2	10	10	8.5	55
	9-3	20	10	9.8	64
	9-4	0	5	7.0	54
	9-5	10	5	6.4	46
	9-6	20	5	6.6	47
	9-7 (control)	0	0	3	11
	9-8 (Gelden®)	>0	--	2	13

25 From these results it is observed that PG has little or no effect on drug delivery at either 5% or 10% of enhancer. However, all the formulations with enhancer provide significantly higher peak and total drug delivery than either the control or the commercial product. There is no 30 significant difference in performance between the control and the commercial product.

Example 10

This example further illustrates the effects of the invention with diclofenac as the NSAID. The test procedure was substantially the same as previously described using either human (H) or porcine (P) skin and an ethanol:water (70:30) vehicle. 2-n-nonyl-1,3-dioxolane was used as the skin permeation enhancer compound according to the invention. The results are shown in Table 12 below. In Run Nos. 10-A through 10-G 1 wt.% of diclofenac (as free acid) was used. In Run Nos. 10-I and 10-J (commercial product) 0.93 wt.% of diclofenac (as free acid) was used.

Table 12

Run No.	Base	PG(%)	Enhancer (%)	Skin	Peak Flux $\mu\text{g}/\text{cm}^2/\text{h}$	% of dose 24h
10-A	Na	20	10	P	12	40
10-B	Na	10	10	P	10	36
10-C	Na	5	10	P	6	30
10-D	Na	20	5	P	8	28
10-E	Na	0	10	P	2.5	19
10-F	Na	20	0	P	1	8
10-G	DEA ^b	20	10	H	15	76
10-H	Na	20	10	H	10	46
10-I	DEA	20	10	H	11	46
10-J ^a	DEA	--	--	H	1.5	10

a - Emugel (Voltaren)

b - diethylamine

From the above results reported in Table 12 the following observations and conclusions may be drawn. The first set of experiments, Run Nos. 10A-10C, show that PG exerts a positive effect as a co-enhancer for diclofenac. In a second set of experiments Run Nos. 10D-10F, it is seen

that the combination of PG with the dioxolane enhancer provides better performance than might be expected from the results with dioxolane enhancer alone and with PG alone. From the third set of experiments, Run Nos. 10-G and 10-H, 5 it is observed that DEA as the counterion (base) provides better performance than sodium (Na). Finally, from the fourth set of experiments, Run Nos. 10-I and 10-J it is seen that the formulation according to the present invention provides significantly improved performance in comparison to 10 a commercial diclofenac topical formulation.

Example 11

This example shows the effects of multiple daily applications of a formulation according to the invention (10% 2-n-nonyl-1,3-dioxolane, 20% PG, 70:30 EtOH/H₂O) and a 15 commercial topical NSAID formulation (Emulgel). Each product was applied to human skin sample with second and third applications following at 8 hour intervals. The results are shown below in Table 13.

Table 13

Run No.	Peak Flux μg/cm ² /h	μg at 24h
11-A (Diclofenac, Na, 0.93%)		
1st Application (0h)	13	125
2nd Application (8h)	11	230
25 3rd Application (16h)	18	360
11-B (Emulgen, Diclofenac-DEA, 0.93%)		
1st Application (0h)	9	100
2nd Application (8h)	4	100
3rd Application (16h)	10	130

These results show that a single application of the topical diclofenac formulation according to the present invention provides comparable performance to 3 applications of the commercial diclofenac topical formulation. Whereas, 5 3 daily applications of the topical NSAID formulation of this invention provides nearly a three-fold higher delivery of drug than 3 applications of the commercial product while a two-fold improvement in drug delivery is obtained with 2 daily applications of the invention product as compared to 2 10 daily applications of the commercial product.

Example 12

Following the same procedures as described in Example 1 and using a standard static cell as described in Example 1, the skin permeation enhancing effect, through human skin, of 15 the compounds of this invention were compared to the skin penetration enhancing effects of other known skin penetration enhancing (SPE) compounds, including, diisopropyl adipate (DIPA), and C₁₂-C₁₅ alkyl benzoate esters (ABE) as well as a mixture of ABE with the invention 20 compound, for a representative NSAID compound, ketoprofen.

Specifically the following compositions (solutions were tested by adding the test compound in the amount shown in the following Table 1 to a vehicle of ethanol/propylene glycol/water at a weight ratio of 70:20:10 and containing 25 2.5% ketoprofen. Each composition, except for the compositions containing DIPA, were tested six times; the DIPA solutions were tested four times. The results of the experiments, in terms of drug delivery ($\mu\text{g}/\text{cm}^2$) at 24 hr and the 24 hour average ($\mu\text{g}/\text{cm}^2/\text{hr}$) are also shown in the 30 following Table 14.

Table 14

Comparison of Delivery of Ketoprofen Through Human Cadaver Skin Using the SPE of this Invention verus other SPE's

<u>Enhancer</u>	<u>Weight%</u>	<u>Drug Delivered ($\mu\text{g}/\text{cm}^2$)</u>	
		<u>24 hours</u>	<u>Average/hr</u>
Control	0	165±114	6.9
2-nonyl-1,3-dioxolane	5	325± 75	13.5
2-nonyl-1,3-dioxolane	10	451± 87	18.8
DIPA	5	166± 66	6.9
DIPA	10	103± 76	4.3
ABE	5	124± 27	5.2
ABE +	5	236± 77	9.8
2-n-nonyl-1,3-dioxolane			

For further clarity the results of the above experiments are graphically represented in the attached Figures 9, 10 and 11 in terms of flux of ketoprofen ($\mu\text{g}/\text{cm}^2/\text{h}$) as a function of time over 24 hours (Fig. 9); cumulative amount of ketoprofen delivered ($\mu\text{g}/\text{cm}^2$) as a function of time (Fig. 10); and cumulative amount of ketoprofen delivered (% of dose) versus time (Fig. 11).

From the results in Table 14 and in Figures 9-11 it is seen that the enhancing effect of the dioxolane, dioxane and acetal compounds of this invention provide significant improvements in enhancement relative to a control with no enhancer whereas the enhancers disclosed in the cited prior art, including diisopropyl adipate and higher fatty alkyl ester of benzoic acid, are no better than the control or

worse than the control. Even when the benzoate enhancer is used in combination with the enhancer of this invention the mixture is less effective than the use of the invention compound alone.

5 Example 13

In a separate set of experiments with 2.5% ketoprofen solutions in the same vehicle as described in Example 12, but using porcine skin, it was observed that similar results were obtained using 10% of decanal dimethyl acetal as the enhancer in place of 10% 2-nonyl-1,3-dioxolane. Also, in the 10% 2-nonyl-1,3-dioxolane containing solution with 2.5% ketoprofen, replacing the propylene glycol component of the vehicle with an equal weight of either glycerol (GL) or propylene carbonate (PC), similar results were obtained.

10 These results are graphically presented in the attached Figures 12, 13 and 14 for flux vs. time, cumulative amount ($\mu\text{g}/\text{cm}^2$) vs. time, and cumulative amount (% of dose) vs. time, respectively.

15 Example 14

20 These experiments were carried out in a similar manner to Example 12 but using porcine skin in place of human skin and gelled compositions rather than solutions. Furthermore, in these experiments the NSAID was ibuprofen at 5%. The enhancers used in this series of experiments included 2-nonyl-1,3-dioxolane (this invention), cineole (eucalyptol) and menthol, each at an amount of 10% by weight of the total composition. In addition to the vehicle (ethanol/propylene glycol/water 70:20:10) the compositions included a gelling agent.

The results are shown in the following Table 15 and in the attached Figures 15, 16 and 17.

Table 15

<u>Enhancer</u>	<u>Amount Ibuprofen Delivered ($\mu\text{g}/\text{cm}^2$)</u>	
	<u>At 24 hours</u>	<u>Average per hour</u>
2-nonyl-1,3-dioxolane	149 ± 26 (1)	6.2
Cineole	36 ± 5.5 (2)	1.5
Menthol	33 ± 7 (1)	1.4

Note (1): based on six replications

(2): based on five replications

These results demonstrate that the cineole and menthol compounds, although disclosed in the art as skin penetration enhancers, are equally ineffective in enhancing the delivery of ibuprofen. In contrast, the SPE compound of this invention is unexpectedly effective in enhancing the transdermal delivery of this NSAID drug.

Example 15

Following the same procedures as described above for Example 12, formulations containing as the NSAID, either the sodium salt of diclofenac or free diclofenac were tested using 10% 2-n-nonyl-1,3-dioxolane, with a carrier (to 100%) comprising ethanol:propylene glycol:water at a weight ratio of 59:20:21. The results are shown in the following Table 16.

Table 16

<u>Drug (salt) (%)</u>	<u>Cumulative Amt. at 24 hours ($\mu\text{g}/\text{cm}^2$)</u>
Na (1)	112
Diethylamine (0.93)	221

This data shows the enhanced benefit of the diethylamine salt over the free acid and demonstrates significant improvement in drug delivery using the inventive SPE compound with the sodium salt of diclofenac.

5 Example 16

The commercial product, Voltaren® Emulgel®, a topical product based on the diethylamine salt of diclofenac, was tested under the same conditions as above. The cumulative amount of drug delivery was only about 7 to 13 $\mu\text{g}/\text{cm}^2/24\text{h}$.

10 When the inventive SPE is used in amounts of 5% or 10%, in a composition similar to Emulgel but with the triethanolamine salt in place of the diethylamine salt, and which includes as the solvent system 20% isopropyl alcohol, 5% propylene glycol, 65.64% water, and 1.2% Carbopol gelling agent, in

15 combination with mineral oil and other emulsifying agents, the 24 hour cumulative delivery is only about 4 to 7 $\mu\text{g}/\text{cm}^2$.

Although not wishing to be bound by any particular theory it is believed that the poor performance in this case is due, not just to the high water content, but to the fact

20 that the SPE is solubilized in the emulsifying agents and is not readily released.

In a separate experiment with diclofenac as free acid, using 10% SPE and an ethanol:propylene glycol:water carrier at a 70:20:10 ratio, the cumulative amount of the free acid

25 which diffused through the human skin sample was as high as about 96 $\mu\text{g}/\text{cm}^2/24\text{h}$.

Example 17

Using the same type of standard static cell as used in Example 1 several different NSAID drugs in the amounts shown in Table 17 using the vehicles and formulation types shown 5 in Table 17 with either human or porcine skin, a series of tests were run to measure the average flux of the formulations with no enhancer or with 5% or 10% of the invention enhancer, 2-n-nonyl-1,3-dioxolane (SEPA®). The results from Table 17 are reported in Table 18 as the 10 Enhancing Factor (ratio of flux for formulation with SEPA® to flux for formulation without SEPA®) to show the degree or extent of improved drug delivery obtained when the invention enhancing compound is added to the formulation.

The data in Tables 17 and 18 clearly demonstrate the 15 effectiveness of the dioxolane, dioxane and acetal enhancers of this invention over a broad range of NSAID compounds in different formulation types.

TABLE 17

AVERAGE FLUX ($\mu\text{g}/\text{cm}^2/\text{h}$)

DRUG	% DRUG	FORMULATION TYPE	SKIN TYPE	AVERAGE 24 Hours FLUX ($\mu\text{g}/\text{cm}^2/\text{h}$)	
Na DICLOFENAC	1%	Solution (70:20:10)	Porcine	No SEPA*	0.98
				5% SEPA*	3.38
				10% SEPA*	4.62
IBUPROFEN	5%	Gel (70:20:10)	Porcine	No SEPA*	19.35
				10% SEPA*	45.5
IBUPROFEN	5%	Gel (70:20:10)	Human	No SEPA*	5.14
				10% SEPA*	15.54
KETOPROFEN	2.5%	Solution (70:20:10)	Porcine	No SEPA*	2.14
				10% SEPA*	7.62
KETOPROFEN	2.5%	Gel (70:20:10)	Porcine	No SEPA*	1.4
				5% SEPA*	5.47
KETOPROFEN	2.5%	Solution (70:30; 20% PG)	Human	10% SEPA*	4.29
				No SEPA*	0.18
KETOPROFEN	2.5%	Solution (70:30; 20% PG)	Human	5% SEPA*	2.18
				10% SEPA*	3.16
KETOROLAC	1%	Gel (70:20:10)	Porcine	No SEPA*	0.99
				10% SEPA*	2.49
KETOROLAC	1%	Gel (70:30; 20% PG)	Porcine	No SEPA*	1.26
				5% SEPA*	3.1
NAPROXEN	5%	Gel (70:30)	Porcine	No SEPA*	0.36
				5% SEPA*	0.94
NAPROXEN	5%	Gel (70:20:10)	Porcine	10% SEPA*	1.14
				No SEPA*	0.17
NAPROXEN	5%	Gel (70:30)	Human	5% SEPA*	1.16
				10% SEPA*	1.55
NAPROXEN	5%	Gel (70:20:10)	Human	No SEPA*	3.38
				5% SEPA*	32.74
NAPROXEN	5%	Gel (70:20:10)	Human	10% SEPA*	24.22
				No SEPA*	1.93
NAPROXEN	5%	Gel (70:20:10)	Human	5% SEPA*	9.81
				10% SEPA*	26.88
PIROXICAM	0.2%	Solution (70:20:10)	Porcine	No SEPA*	0.01
				10% SEPA*	0.08
PIROXICAM	0.5%	Solution (70:20:10)	Porcine	No SEPA*	0.21
				5% SEPA*	0.99
PIROXICAM	0.5%	Gel (70:20:10)	Porcine	10% SEPA*	1.21
				No SEPA*	0.09
PIROXICAM	0.5%	Gel (70:30)	Human	5% SEPA*	0.44
				10% SEPA*	0.42
PIROXICAM	0.5%	Gel (70:30)	Human	No SEPA*	0.76
				5% SEPA*	3.53
PIROXICAM	0.5%	Gel (70:30)	Human	10% SEPA*	4.36

TABLE 18

SEPA/NSAID ENHANCING FACTOR

DRUG	% DRUG	FORMULATION TYPE	SKIN TYPE	ENHANCING FACTOR	
Na DICLOFENAC	1%	Solution (70:20:10)	Porcine	No SEPA* vs 5% SEPA*	3.5
				No SEPA* vs 10% SEPA*	4.7
IBUPROFEN	5%	Gel (70:20:10)	Porcine	No SEPA* vs 10% SEPA*	3
		Gel (70:20:10)	Human	No SEPA* vs 10% SEPA*	2.7
KETOPROFEN	2.5%	Solution (70:20:10)	Porcine	No SEPA* vs 10% SEPA*	3.8
				No SEPA* vs 5% SEPA*	3.9
KETOPROFEN	2.5%	Gel (70:20:10)	Porcine	No SEPA* vs 10% SEPA*	3.1
				No SEPA* vs 5% SEPA*	12.8
KETOPROFEN	2.5%	Solution (70:30; 20% PG)	Human	No SEPA* vs 10% SEPA*	18.6
				No SEPA* vs 5% SEPA*	
KETOROLAC	1%	Gel (70:20:10)	Porcine	No SEPA* vs 10% SEPA*	7.7
		Gel (70:30; 20% PG)	Porcine	No SEPA* vs 5% SEPA*	2.5
				No SEPA* vs 10% SEPA*	2.1
NAPROXEN	5%	Gel (70:30)	Porcine	No SEPA* vs 5% SEPA*	2.7
				No SEPA* vs 10% SEPA*	3.1
NAPROXEN	5%	Gel (70:20:10)	Porcine	No SEPA* vs 5% SEPA*	7
				No SEPA* vs 10% SEPA*	8.9
NAPROXEN	5%	Gel (70:30)	Human	No SEPA* vs 5% SEPA*	10
				No SEPA* vs 10% SEPA*	6.9
NAPROXEN	5%	Gel (70:20:10)	Human	No SEPA* vs 5% SEPA*	5.2
				No SEPA* vs 10% SEPA*	13.6
PIROXICAM	0.2%	Solution (70:20:10)	Porcine	No SEPA* vs 10% SEPA*	6.4
				No SEPA* vs 5% SEPA*	5
PIROXICAM	0.5%	Solution (70:20:10)	Porcine	No SEPA* vs 10% SEPA*	6.2
				No SEPA* vs 5% SEPA*	5.9
PIROXICAM	0.5%	Gel (70:20:10)	Porcine	No SEPA* vs 10% SEPA*	6
				No SEPA* vs 5% SEPA*	4.8
PIROXICAM	0.5%	Gel (70:30)	Human	No SEPA* vs 10% SEPA*	5.9

What is claimed is:

1 Claim 1. A substantially neutral ibuprofen containing
2 alcoholic or aqueous alcoholic composition which comprises,
3 on a weight basis, of the total composition:

4 a therapeutically effective amount of ibuprofen in the
5 form of a pharmacologically acceptable salt of;

6 a skin penetration enhancing effective amount of a C₇
7 to C₁₄-hydrocarbyl substituted 1,3-dioxolane, 1,3-dioxane or
8 acetal;

9 0 to about 18% of glycol having from 3 to 6 carbon
10 atoms;

11 at least about 40% of volatile alcohol selected from
12 the group consisting of ethanol, propanol and mixture
13 thereof;

14 0 to about 40% water;

15 base to provide a pH in the range of from about 6 to
16 about 8, and,

17 optionally, a gelling agent effective to thicken the
18 composition to avoid or minimize run-off when applied to the
19 skin.

1 Claim 2. The composition according to Claim 1 which
2 comprises

3 from about 2 to 10% ibuprofen;

4 from about 4 to 15% of the enhancer wherein the
5 hydrocarbyl group substituent has from about 7 to 10 carbon
6 atoms;

7 from about 0 to 15% propylene glycol;

8 from about 55 to 70% ethanol;
9 from about 4 to 35% water;
10 base in amount to adjust the pH of the composition in
11 the range of from about 6.5 to about 7.5, and,
12 0 to about 2% of cellulosic thickener.

1 Claim 3. The composition according to claim 2 which
2 comprises

3 from about 2 to 10% ibuprofen;
4 from about 5 to 10% of the enhancer;
5 about 0% propylene glycol;
6 from about 55 to 70% ethanol;
7 from about 4 to 35% water;
8 based in amount to adjust the pH of the composition to
9 from about 6.5 to about 7.5; and
10 0 to 2% gelling agent.

1 Claim 4. The composition according to claim 2 which
2 comprises

3 from about 2 to 10% ibuprofen;
4 from about 5 to 10% of the enhancer;
5 from about 1 to 15% propylene glycol;
6 from about 55 to 70% ethanol;
7 from about 4 to 35% water;
8 base in amount to adjust the pH of the composition to
9 from about 6.5 to about 7.5; and
10 0 to 2% gelling agent.

1 Claim 5. The composition of claim 1 which comprises
2 about 5% ibuprofen;

3 from about 5 to 10% skin penetration enhancer wherein
4 the hydrocarbyl group substituent has from about 7 to 10
5 carbon atoms;

6 up to about 5% propylene glycol;
7 from about 55 to 70% ethanol;
8 water in amount to provide an ethanol:water ratio, by
9 weight, of about 70:30;
10 base in amount to substantially neutralize the
11 ibuprofen;
12 and, gelling agent in amount effective to thicken the
13 composition.

1 Claim 6. The composition of claim 5 which is free from
2 propylene glycol.

1 Claim 7. The composition of claim 1 which comprises
2 2-n-nonyl-1,3-dioxolane as skin penetration enhancing
3 compound.

1 Claim 8. A substantially neutral alcoholic or aqueous
2 alcoholic topical composition effective for the transdermal
3 delivery of non-steroidal anti-inflammatory drug which
4 comprises, based on the weight of the total composition,
5 a therapeutically effective amount of a
6 pharmacologically acceptable salt of a non-steroidal
7 antiinflammatory drug selected from the group consisting of
8 tolmetin, diclofenac, keterolac, arylpropionic acids (other
9 than ibuprofen), anthranilic acids, enolic acids, alkanones,
10 sulindac and etodolac;
11 from about 0.5 to 25 % of C₆ to C₁₄-hydrocarbyl
12 derivative of 1,3-dioxolane, 1,3-dioxane or acetal as skin
13 penetrant enhancer;
14 0 to about 40% of glycol having from 3 to 6 carbon
15 atoms;

16 at least about 40% of volatile alcohol selected from
17 the group consisting of ethanol, propanol and mixtures
18 thereof;

19 up to about 40% water;

20 base in an amount to provide a pH of from about 6 to
21 about 8, and

22 up to about 5% gelling agent.

1 Claim 9. The composition of claim 3 which comprises
2 2-n-nonyl-1,3-dioxolane as skin penetration enhancing
3 compound.

1 Claim 10. The composition according to claim 8 which
2 comprises:

3 from about 0.1 to 10% diclofenac, ketorolac, naproxen,
4 flurbiprofen, ketoprofen or piroxicam;

5 from about 2 to 15% of the skin penetration enhancer;

6 0 to about 30% propylene glycol;

7 from about 45 to 70% ethanol, isopropanol or mixture
8 thereof;

9 0 to about 20% water;

10 from about 0.1 to 3% gelling agent, and

11 base to provide a pH in the range of from about 6.5 to
12 about 7.5.

1 Claim 11. The composition of claim 4 wherein the non-
2 steroidal antiinflammatory is diclofenac or ketoprofen.

1 Claim 12. The composition of claim 11 wherein the
2 amount of propylene glycol is in the range of from about 5
3 to 20%.

1 Claim 13. The composition of claim 12 wherein the
2 amount of skin penetration enhancer is from about 5 to about
3 10%.

1 Claim 14. The composition of claim 4 wherein the non-
2 steroidal antiinflammatory drug is diethanloamine salt of
3 diclofenac.

1 Claim 15. A glycol-free topical composition effective
2 for the transdermal administration of naproxen, which
3 comprise, on a weight basis of the total composition:

4 a pharmaceutically effective amount of naproxen,
5 from about 2 to 20% of 2-C₇-C₁₄ hydrocarbyl substituted
6 1,3-dioxolane, 1,3-dioxane, or acetal skin penetration
7 enhancer;

8 from about 50 to 85% ethanol, iso-propanol, or mixture
9 thereof;

10 0 to about 40% water;

11 base in an amount to provide a pH in the range of from
12 about 6 to about 8, and

13 up to about 5% gelling agent.

1 Claim 16. The glycol-free composition of claim 5
2 wherein the amount of enhancer is from about 5 to 10%.

1 Claim 17. The composition of claim 4 wherein the non-
2 steroidal antiinflammatory drug is diethanloamine salt of
3 diclofenac.

1 Claim 18. A method for the transdermal administration
2 of ibuprofen to a patient in need thereof which comprises
3 topically applying to the skin of the patient a
4 substantially neutral composition comprising from about 5 to
5 15 weight percent of ibuprofen in a vehicle comprising a
6 lower alcohol selected from the group consisting of ethanol,
7 isopropanol and mixture thereof, alkyl glycol having from 3
8 to 6 carbon atoms, and water in a mixing ratio of
9 alcohol:glycol:water of 40-80:0-20:0-40, said vehicle

10 comprising from about 70 to 90 weight percent of the
11 composition, and from about 5 to 15 weight percent of a skin
12 penetration enhancing compound selected from the group
13 consisting of 2-hydrocarbyl-1,3-dioxolane, 2-hydrocarbyl-
14 1,3-dioxane and hydrocarbyl substituted-acetal, wherein the
15 hydrocarbyl group has from 7 to 14 carbon atoms.

1 Claim 19. A method for the transdermal administration
2 of a non-steroidal antiinflammatory drug selected from the
3 group consisting of tolmetin, diclofenac, keterolac,
4 arylpropionic acids (other than ibuprofen), anthranilic
5 acids, enolic acids, alkanones, sulindac and etodolac to a
6 patient in need thereof which comprises

7 topically applying to the skin of the patient a
8 substantially neutral composition comprising from about 0.1
9 to 10 weight percent of the non-steroidal antiinflammatory
10 drug in a vehicle comprising a lower alcohol selected from
11 the group consisting of ethanol, isopropanol and mixture
12 thereof, alkyl glycol having from 3 to 6 carbon atoms, and
13 water in a mixing ratio of alcohol:glycol:water of 40-80:0-
14 40:0-40, said vehicle comprising from about 70 to 90 weight
15 percent of the composition, and from about 0.5 to 25 weight
16 percent of a skin penetration enhancing compound selected
17 from the group consisting of 2-hydrocarbyl-1,3-dioxolane, 2-
18 hydrocarbyl-1,3-dioxane and hydrocarbyl substituted-acetal,
19 wherein the hydrocarbyl group has from 7 to 14 carbon atoms.

1 Claim 20. A method for the transdermal administration
2 of naproxen to a patient in need thereof which comprises
3 topically applying to the skin of the patient a
4 substantially neutral composition comprising a
5 therapeutically effective amount of naproxen in a glycol-

6 free vehicle comprising a lower alcohol selected from the
7 group consisting of ethanol, isopropanol and mixture
8 thereof, and water in a mixing ratio of alcohol:water of 50-
9 85:10-40, said vehicle comprising from about 70 to 90 weight
10 percent of the composition, and from about 2 to 20 weight
11 percent of a skin penetration enhancing compound selected
12 from the group consisting of 2-hydrocarbyl-1,3-dioxolane, 2-
13 hydrocarbyl-1,3-dioxane and hydrocarbyl substituted-acetal,
14 wherein the hydrocarbyl has from 7 to 14 carbon atoms.

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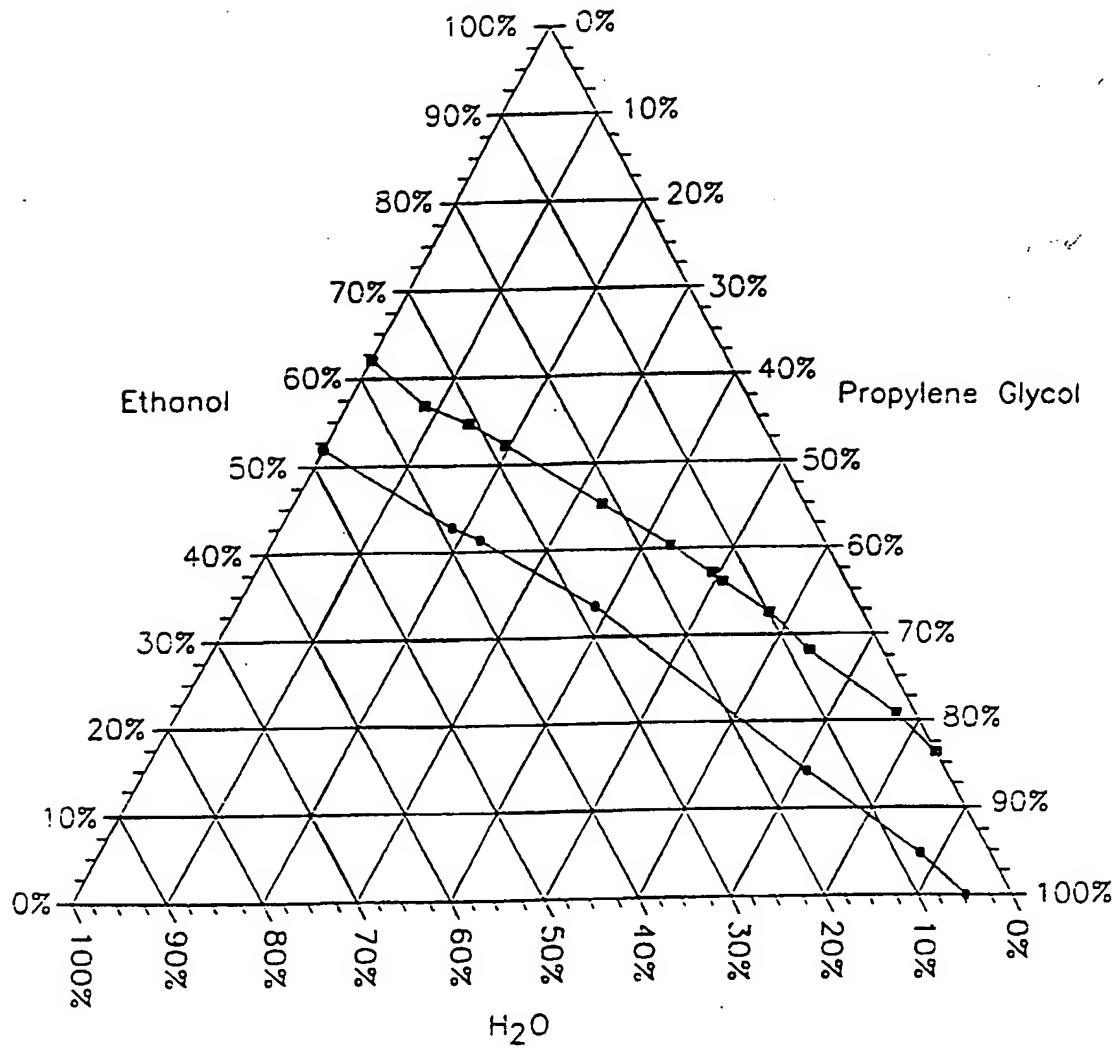


FIG. I

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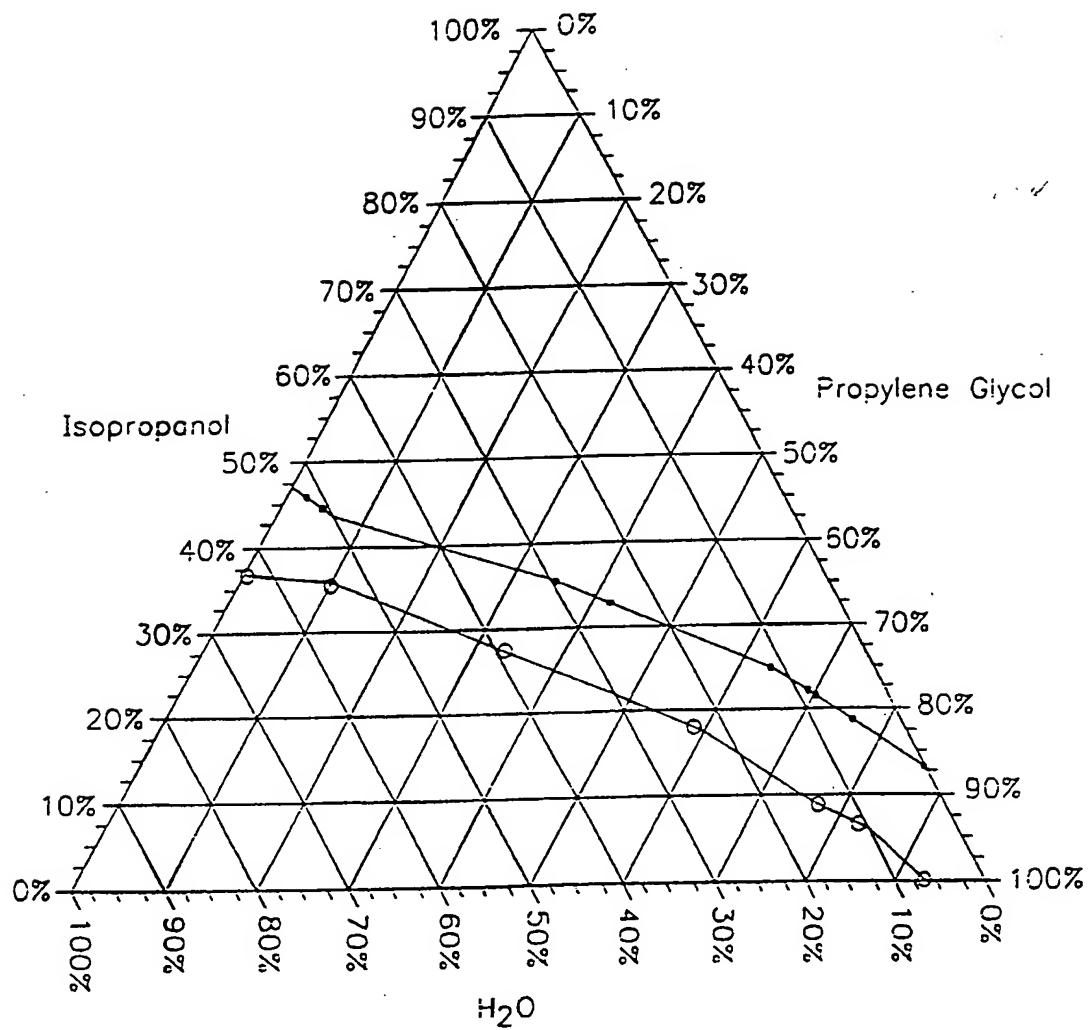


FIG. 2

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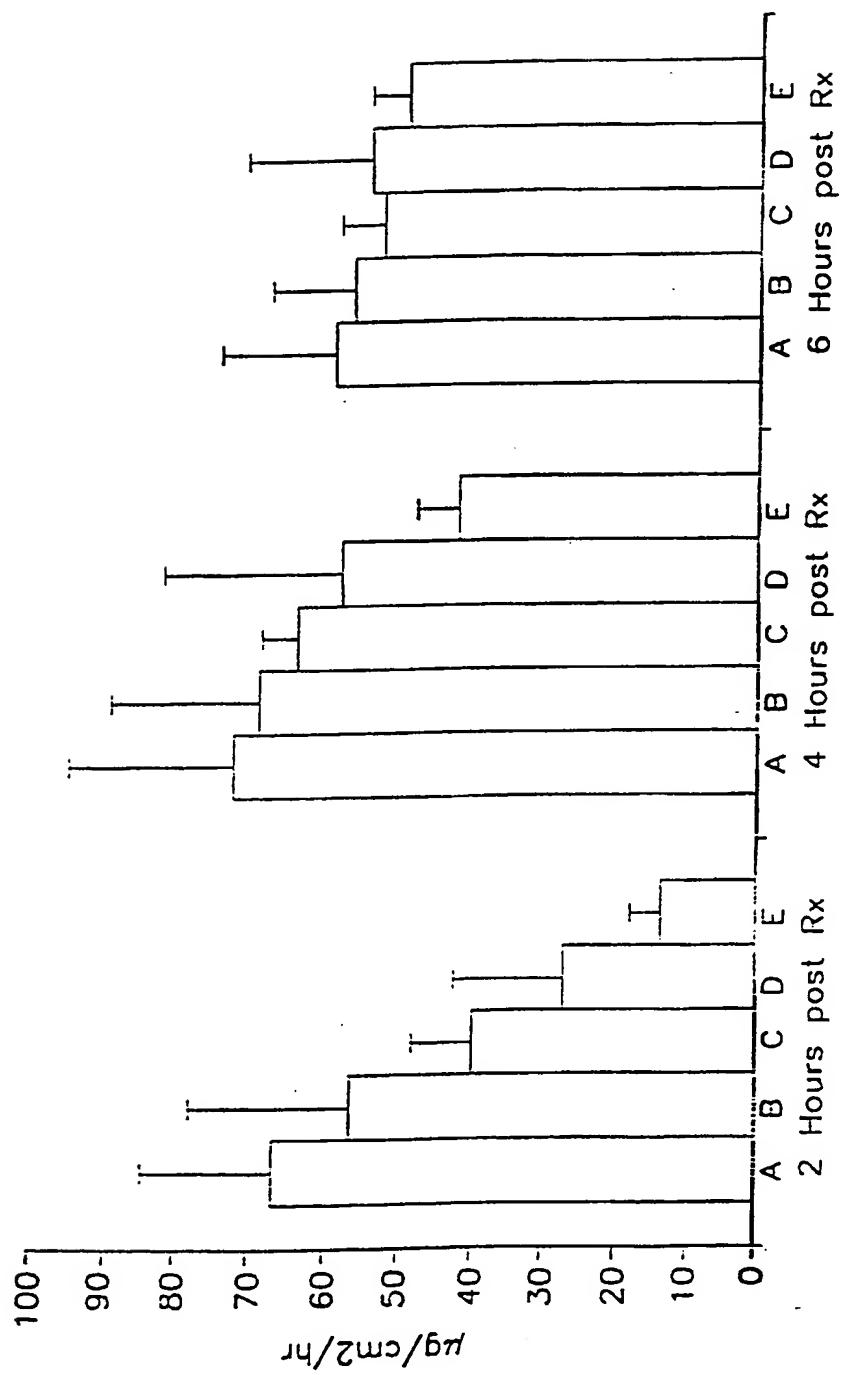
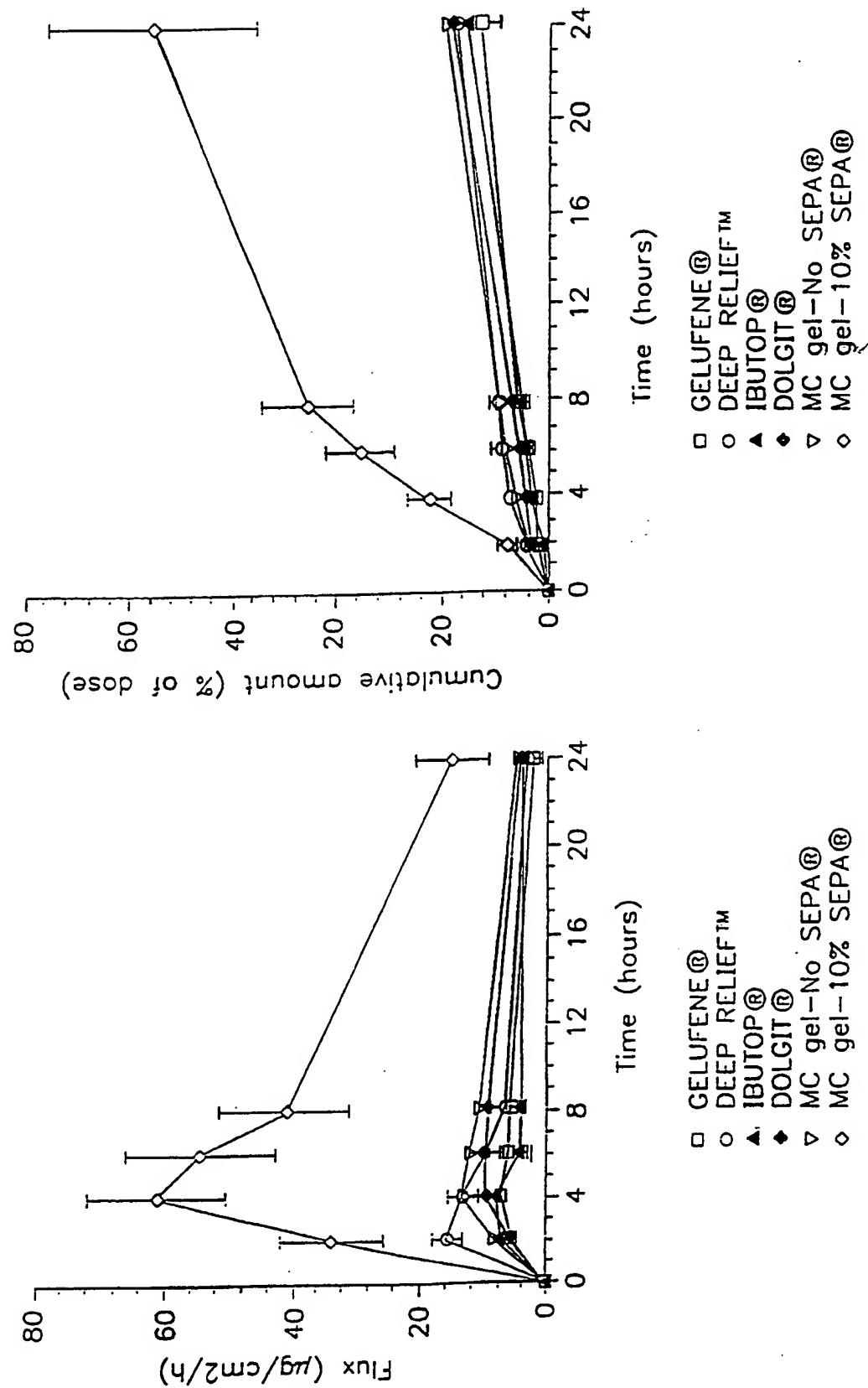


FIG. 3

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**FIG. 5**

5 / 15

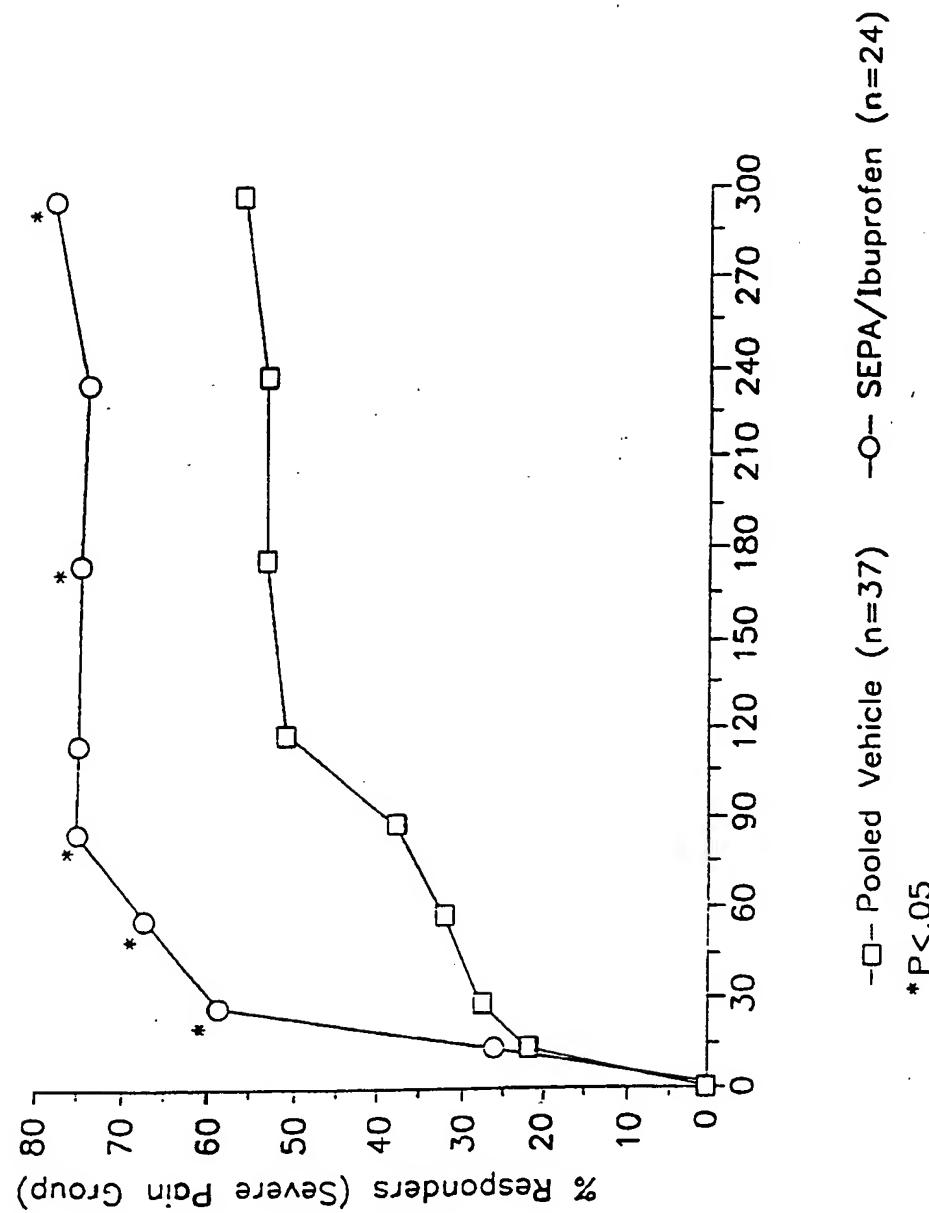


FIG. 6

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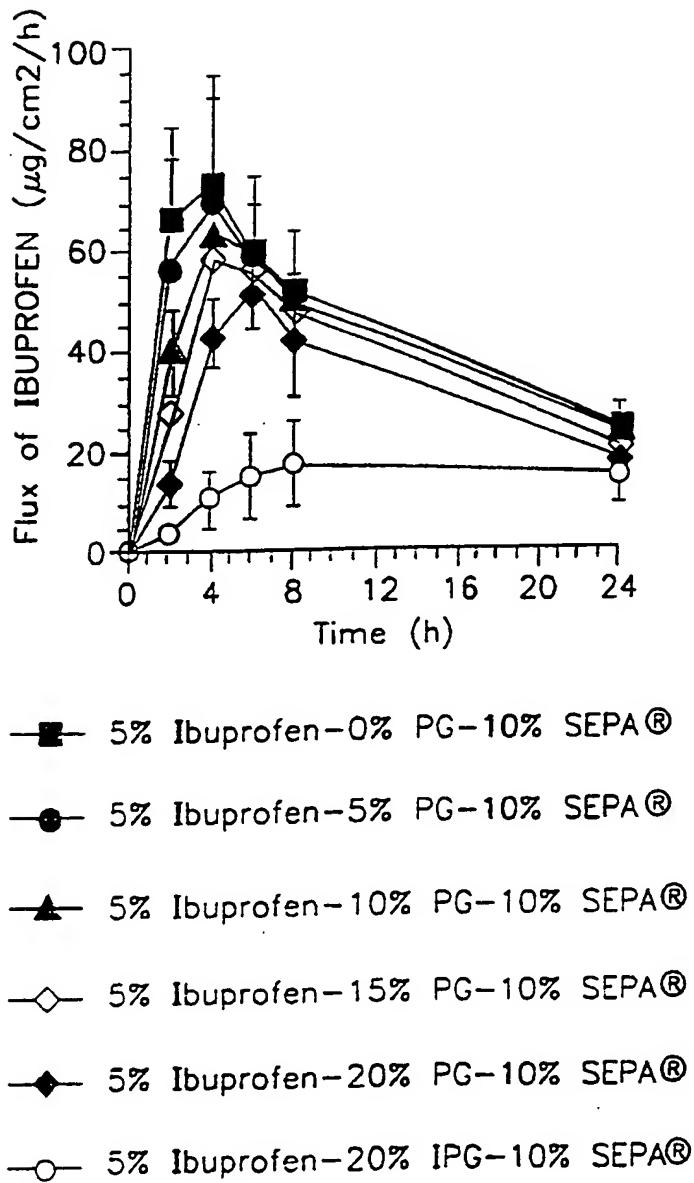
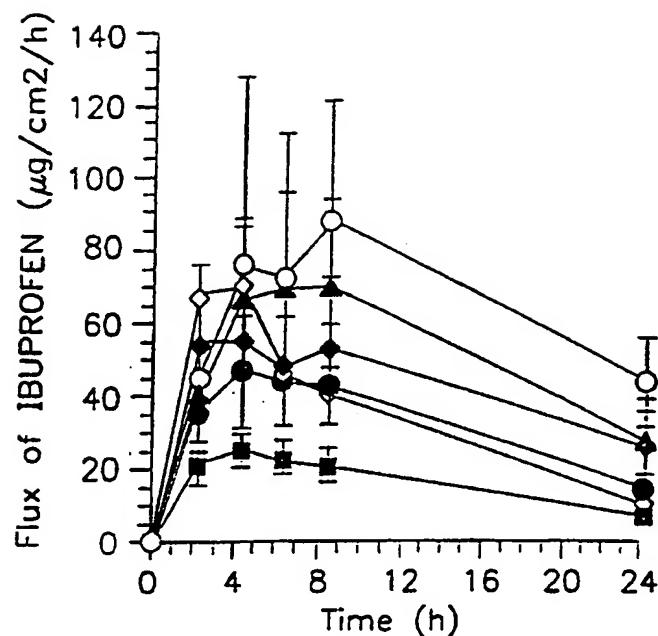


FIG. 7

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- 2.5% Ibuprofen gel (E:PG:W:70:20:10)-10% SEPA®
- 5% Ibuprofen gel (E:PG:W:70:20:10)-10% SEPA®
- ▲ 10% Ibuprofen gel (E:PG:W:70:20:10)-10% SEPA®
- ◇ 2.5% Ibuprofen gel (E:W:70:30)-10% SEPA®
- ◆ 5% Ibuprofen gel (E:W:70:30)-10% SEPA®
- 10% Ibuprofen gel (E:W:70:30)-10% SEPA®

FIG. 8

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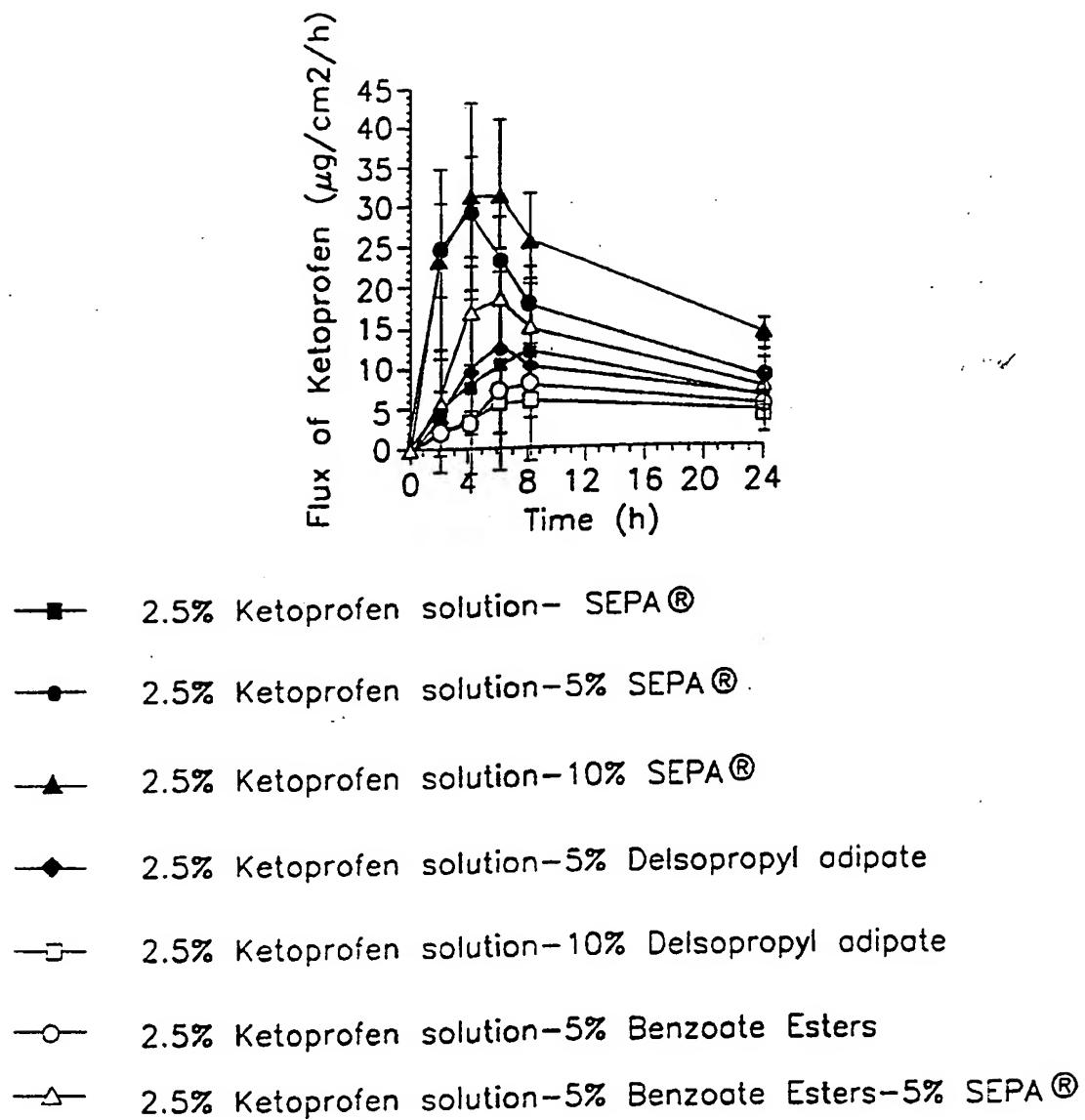


FIG. 9

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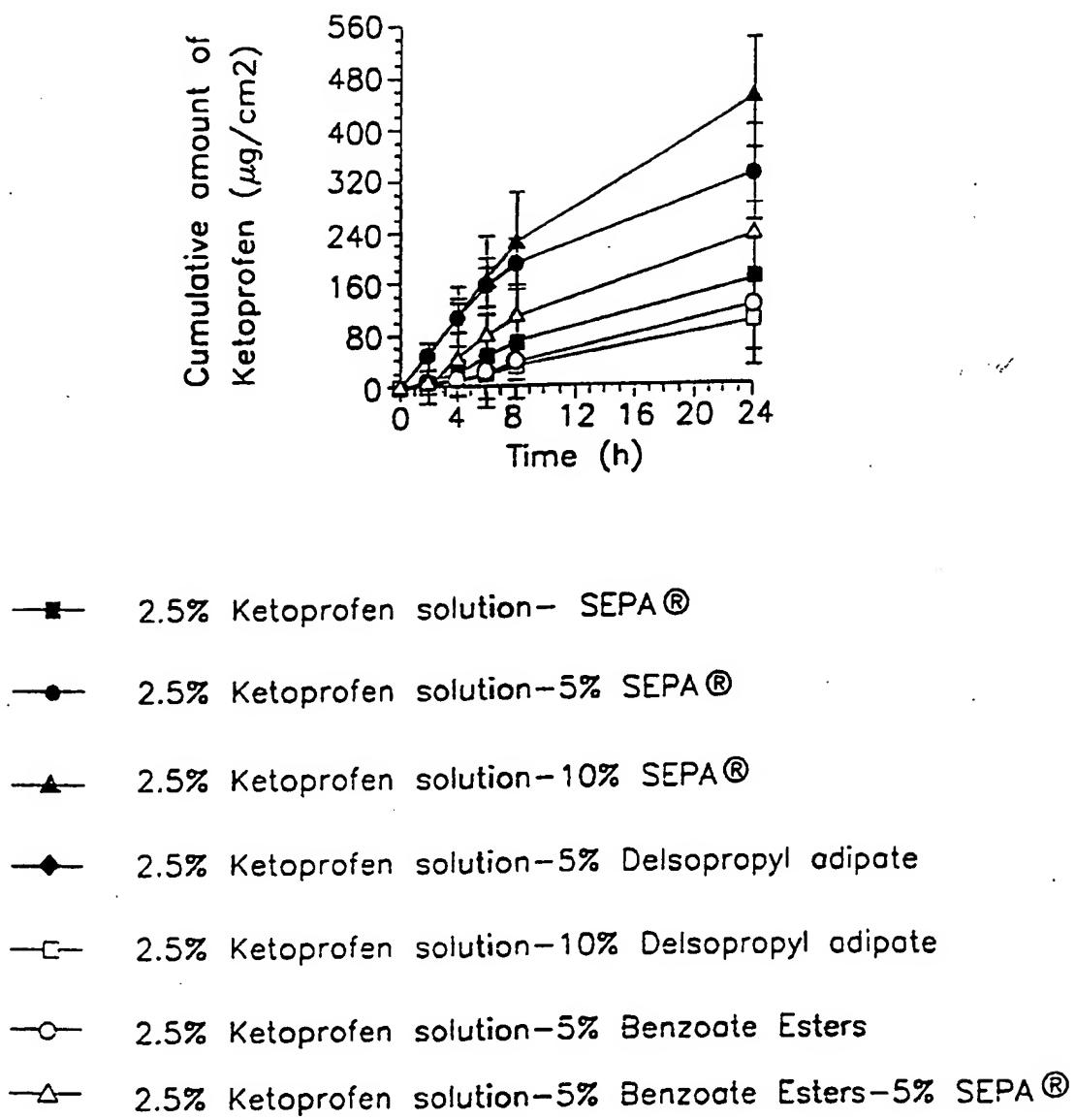
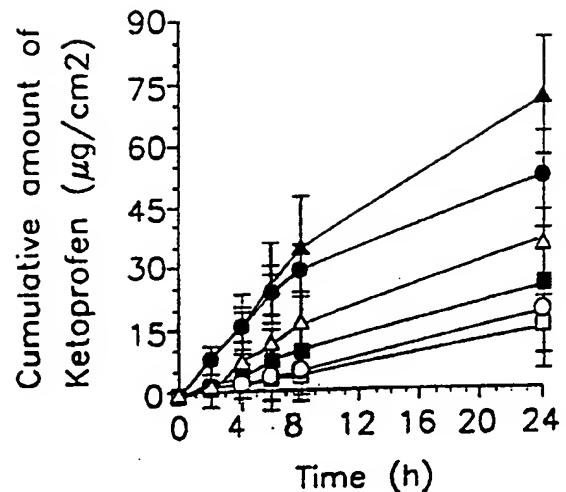


FIG. 10

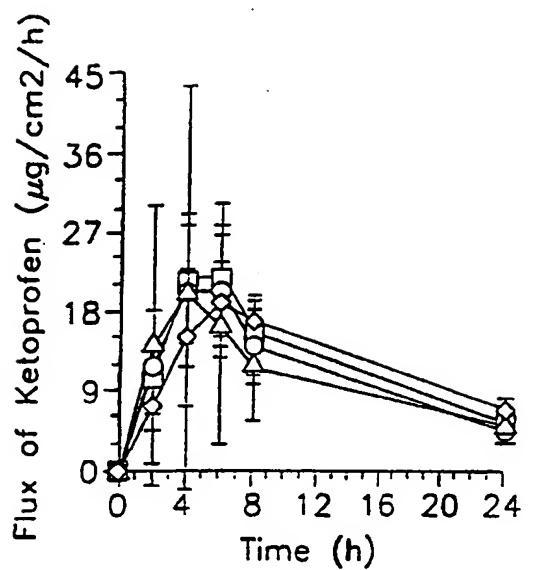
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- 2.5% Ketoprofen solution- SEPA®
- 2.5% Ketoprofen solution-5% SEPA®
- ▲ 2.5% Ketoprofen solution-10% SEPA®
- ◆ 2.5% Ketoprofen solution-5% Delsopropyl adipate
- 2.5% Ketoprofen solution-10% Delsopropyl adipate
- 2.5% Ketoprofen solution-5% Benzoate Esters
- △ 2.5% Ketoprofen solution-5% Benzoate Esters-5% SEPA®

FIG. II

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- 2.5% Ketoprofen solution (E:PG:W; 70:20:10)-10% SEPA®
- 2.5% Ketoprofen solution (E:PG:W; 70:20:10)-10% Decanal DMA
- △— 2.5% Ketoprofen solution (E:GL:W; 70:20:10)-10% SEPA®
- ◇— 2.5% Ketoprofen solution (E:PC:W; 70:20:10)-10% SEPA®

FIG. 12

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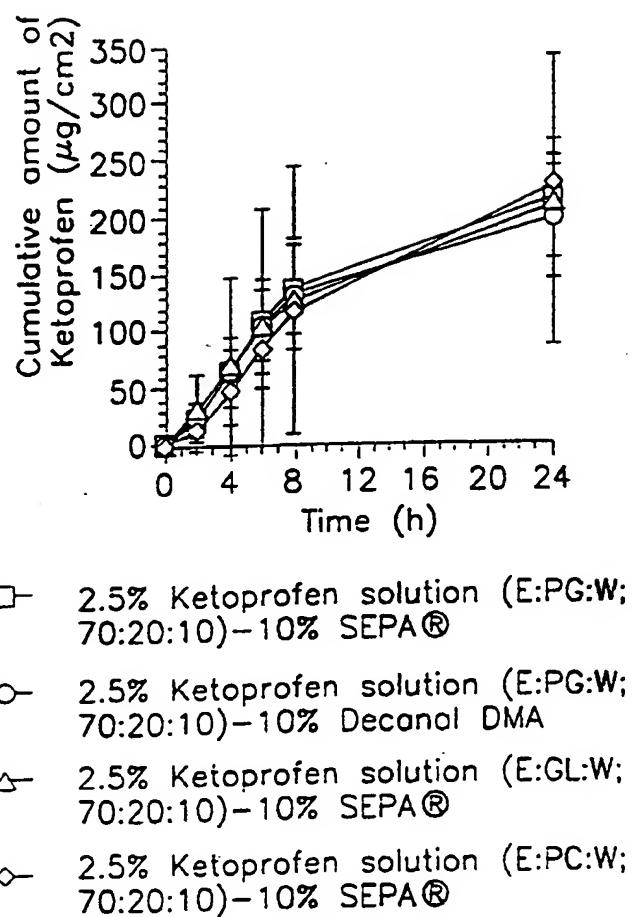


FIG. 13

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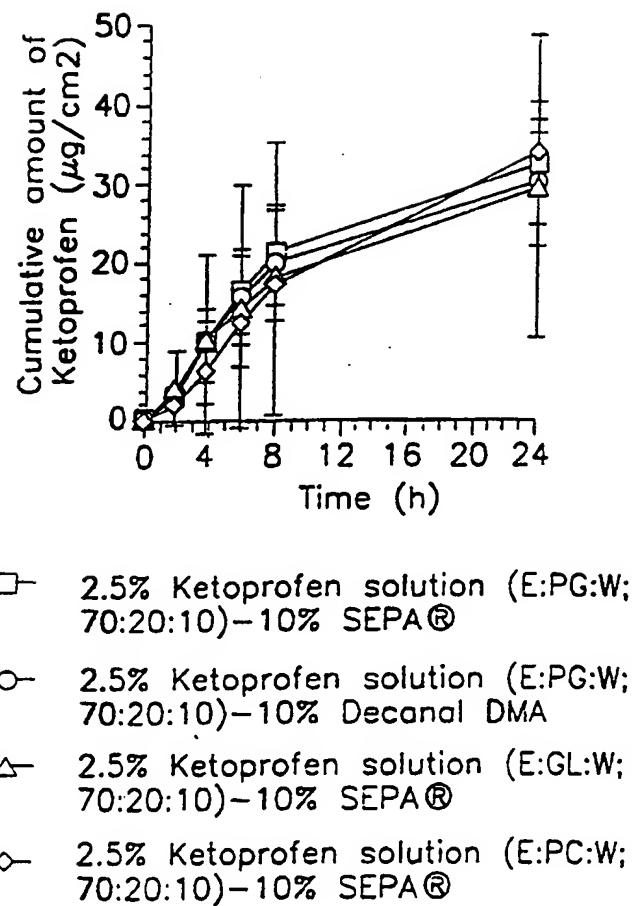
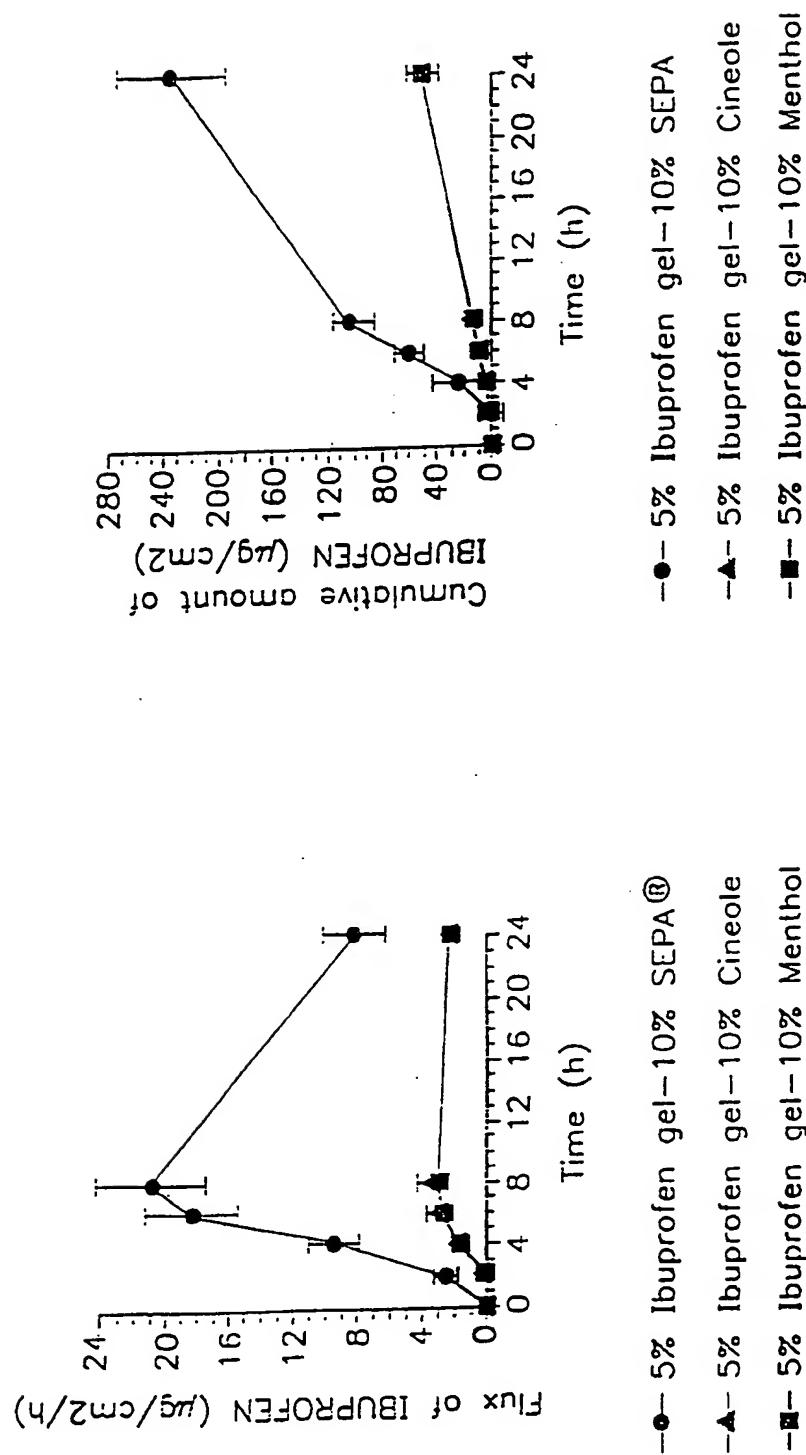


FIG. 14

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**FIG. 16**

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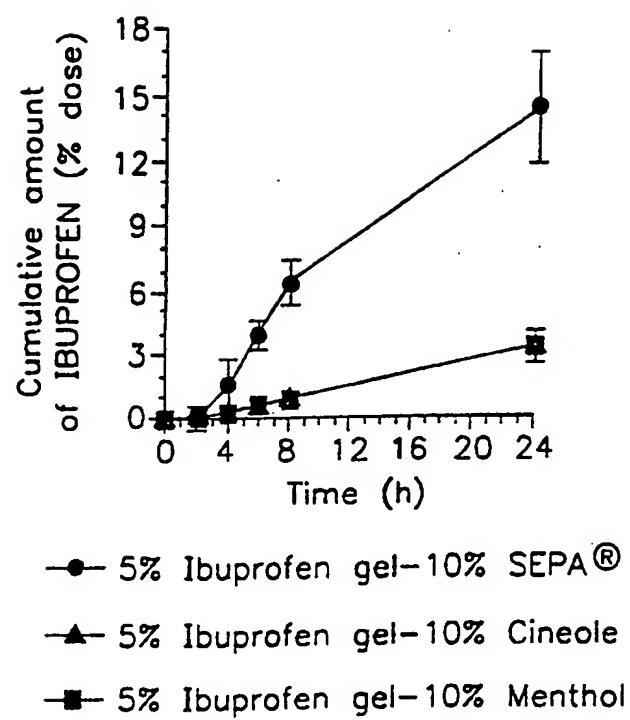


FIG. 17

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US98/17523

A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) :A61K 9/10, 47/24
US CL :424/484, 449; 514/944, 946

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 424/484, 449; 514/944, 946

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	US 4,861,764 A (SAMOUR et al.) 29 August 1989, abstract.	1-14, 18, 19
Y	US 5,093,133 A (WISNIEWSKI et al.) 03 March 1992, abstract.	1-7, 9, 18
Y	US 4,185,100 A (ROVEE et al.) 22 January 1980, col.6, lines 9-12, col. 3, lines 3-8.	1-13, 18, 19.
Y	US 4,533,546 A (KISHI et al.) 06 August 1985, abstract, col. 2 line 57-col. 3, lines 44-47.	1, 2, 5-13, 18, 19
Y	US 4,393,076 A (NODA et al.) 12 July 1983, abstract, col. 3, lines 16-20.	8-13, 19

Further documents are listed in the continuation of Box C.

See patent family annex.

"	Special categories of cited documents:	"T"	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A"	document defining the general state of the art which is not considered to be of particular relevance	"X"	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"E"	earlier document published on or after the international filing date	"Y"	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"L"	document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"&"	document member of the same patent family
"O"	document referring to an oral disclosure, use, exhibition or other means		
"P"	document published prior to the international filing date but later than the priority date claimed		

Date of the actual completion of the international search

17 NOVEMBER 1998

Date of mailing of the international search report

23 DEC 1998

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